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73 Patentinhaber:

PGI Graphics Imaging LLC, Waltham, Mass., US

74 Vertreter:

Patent- und Rechtsanwälte Kraus & Weisert, 80539
München

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72 Erfinder:

ELLIS, W., Ernest, Leverett, US; FOLEY, M., Diane,
Northampton, US; ARNOLD, R., Dana,
Northampton, US

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(54) **IMPROVED ABLATION-TRANSFER IMAGING/RECORDING**

BILDERZEUGUNG/AUFZEICHNUNG DURCH VERBESSERTE ABTRAGUNGS-ÜBERTRAGUNG
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(73) Proprietor: **PGI Graphics Imaging LLC**
Waltham, Mass. 02154 (US)

(72) Inventors:
• **ELLIS, Ernest, W.**
Leverett, MA 01054 (US)
• **FOLEY, Diane, M.**
Northampton, MA 02158 (US)
• **ARNOLD, Dana, R.**
Northampton, MA 02165 (US)

(74) Representative: **Kraus, Walter, Dr. et al**
Patentanwälte Kraus, Welsert & Partner
Thomas-Wimmer-Ring 15
80539 München (DE)

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US-A- 4 702 958	US-A- 4 711 834
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- **PATENT ABSTRACTS OF JAPAN vol. 12, no. 346**
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Description

[0001] The present invention relates a method for transferring a contrasting pattern of intelligence from an ablation-transfer imaging media to a receptor element in contiguous registration therewith, and to an ablation-transfer imaging medium for use in this process.

[0002] The phenomenon of, e.g., laser-induced ablation-transfer imaging, is generically known to this art and is believed to entail both complex non-equilibrium physical and chemical mechanisms. Indeed, such laser-induced ablation-transfer is thought to be effected by the rapid and transient accumulation of pressure beneath and/or within a mass transfer layer initiated by imagewise irradiation. Transient pressure accumulation can be attributed to one or more of the following factors: rapid gas formation via chemical decomposition and/or rapid heating of trapped gases, evaporation, photo and thermal expansion, ionization and/or by propagation of a shockwave. The force produced by the release of such pressure is sufficient to cause transfer of the imaging layer to an adjacent receptor element. The force is preferably sufficient to effect the complete transfer of the exposed area of an entire layer rather than the partial or selective transfer of components thereof.

[0003] Other material transfer imaging/recording techniques based on equilibrium physical changes in the material are also known to this art, but are severely limited in terms of both the overall speed of the process as well as in the materials which can be employed therefor. In particular, ablation transfer differs from the known material transfer techniques such as, for example, thermal melt transfer and dye sublimation/dye diffusion thermal transfer (D2T2). Each of these prior art techniques typically employs thermal print heads as the source of imaging energy.

[0004] Alternatively, it is known to employ laser heating in lieu of the thermal printing head. In these systems, the donor sheet includes a material which strongly absorbs at the wavelength of the laser emission. In the thermal melt transfer process, when the donor sheet is irradiated, this absorbing material converts the laser light to thermal energy and transfers the heat to a colorant transfer layer which also includes a binder, fusible compound, etc., thereby raising its temperature above its melting point to effect its transfer onto an adjacent receptor sheet. In the D2T2 process, only the colorant is transferred to a specially treated or special receptor sheet (e.g., coated or porous) by sublimation or thermal diffusion. See, for example, JP-A-62-140 884, GB-A-2 083 726, US-A-4 804 975, US-A-4 804 977, US-A-4 876 235, US-A-4 753 923 and US-A-4 912 083.

[0005] Compare also US-A-3 745 586 relating to the use of laser energy to selectively irradiate the uncoated surface of a thin film, coated on one side with a contrast imaging absorber, to vaporize and to cause the selective transfer of the absorber coating to an adjacently spaced receptor, and US-A-3 978 247 relating to sublimation transfer recording via laser energy (laser addressed D2T2), wherein the contrast imaging material is also the absorber.

[0006] The aforementioned JP-A-62-140 884 describes a laser recording device, the focus of which is a composite transfer sheet including a base layer of plastic film, a middle layer containing substances that absorb laser light and convert it to heat and a transfer layer topcoat, the primary component of which is a colorant, as well as a receiver sheet to which images are transferred under the influence of the laser. The colorant transfer layer and/or top receiver sheet layer necessarily includes a heat-fusible phenolic, naphtholic, aromatic carboxylic acid or fatty acid amide compound. The plastic film base layer is thus coated with a substance that absorbs laser light and converts it to heat and is not transferred. This heat melts and transfers the image from the transfer layer to the receiver sheet and is said to provide a high-resolution and high-density image thereon.

[0007] Other transfer-imaging systems are described in US-A-4 123 578, US-A-4 157 412, US-A-4 554 238, US-A-4 847 237 and US-A-4 897 310.

[0008] Nonetheless, these processes are limited in a variety of significant respects. For example, in melt transfer, the composition must contain low melting materials to transfer a pigment or dye and receptor sheets appropriately textured for wicking or having special coatings are required for best results. In D2T2, only the imaging dye itself is transferred; thus, it becomes necessary to employ special receptor sheets in order to effectively bind and stabilize ("trap") the dye. Compare, for example, US-A-4 914 078. Furthermore, additional post-heating treatment steps, such as the "setting" of the dyes in the binder which is present on the receptor sheet increases both the complexity and the time associated with the process. Such process is also limited to those dyes and pigments which undergo sublimation or diffusion in response to the particular imaging stimulus.

[0009] These processes are further limited in that the relatively slow processes of heat diffusion and thermal equilibrium are involved.

[0010] Accordingly, need exists in this art for a transfer process which is far more rapid than current transfer techniques, which can effectively employ a wide variety of functional materials and which is not limited to specially treated or special receptor elements.

[0011] Laser-induced recording based on the removal or displacement of material from the exposed area is also known to the recording art. However, these applications do not require transfer of the material from one substrate to another. Historically, laser-induced recording has been used, for example, in optical disk writing with near infrared (IR) lasers typically emitting at wavelengths ranging from 760 nm to 850 nm employed as the writing source. Since polymeric

binders are typically non-absorbent in the near infrared region (760 nm to 2500 nm), infrared absorbers, i.e., sensitizers, are added to the substrate to absorb the laser radiation. This arrangement allows the laser radiation absorbed by the sensitizer to be converted to heat which causes pit formation. See, for example, US-A-4 415 621, US-A-4 446 233, US-A-4 582 776, US-A-4 809 022 and N. Shimadzu et al., The Journal of Imaging Technology, Vol. 15, No. 1, pg. 19 (1989). However, because this technology does not entail the imagewise transfer of materials from one substrate to another, these systems will not be further discussed.

[0012] Prior art laser-induced ablative transfer imaging processes are limited to the use of large amounts of a black body absorber such as graphite or carbon black to transfer a black image. See, for example, US-A-4 245 003, US-A-4 702 958 and US-A-4 711 834 (graphite sensitizer/absorber), US-A-4 588 674 (carbon black sensitizer/absorber), and GB-A-2 176 018 (small amounts of Cyasorb IR 165, 126 or 99 infrared absorbers in combination with graphite as the sensitizer/absorber). Since the black body absorbers employed are highly absorbent in the visible and ultraviolet (UV) as well as in the infrared region, the resulting transferred image is always black due to the presence of the absorber. Such ablative transfer imaging based on black body absorbers is therefore entirely ineffective and wholly unsuited for many applications, e.g., color transfer imaging, color proofing, security printing, etc.

[0013] US-A-3 962 513 describes a laser-ablative transfer medium intended for use in producing a planographic printing plate. This medium has a laser energy absorbing layer, which comprises particles of carbon black and a self-oxidizing binder, and which is separate from a topcoat comprising an ink-receptive resin. Upon absorption of laser radiation by the energy absorbing layer, the topcoat is ablatively transferred to a lithographic surface. Inevitably, at least a portion of the energy absorbing layer will be transferred with the topcoat; the patent states:

[I]t is believed that the products of combustion or blow-off from the irradiated area of the laser responsive layer are absorbed in the portion of the ink-receptive layer which has been transferred to the lithographic surface, thereby limiting scattering or dispersal of such products, which would cause a diffuse image.

Thus, a black image will be created by the carbon black transferred from the energy absorbing layer. Such a black image is of course not objectionable in a printing plate, since only the ink absorbed by the ink-receptive resin is ultimately transferred to the printed stock.

[0014] Thus, a serious need continues to exist for a cost effective ablative transfer imaging medium that is sensitized independently of the contrast imaging material(s), e.g., colorants, and which, therefore, was not limited to contrast materials which must absorb the imaging radiation.

[0015] This invention provides an ablation-transfer imaging medium, and a process for its use, which is similar to that described in the aforementioned US-A-3 962 513 in that it comprises a radiation-absorbing "dynamic release layer" which is separate from a topcoat containing a colorant. However, in the present medium and process, the dynamic release layer uses a non-black body radiation absorber, and the topcoat contains a non-black body contrast imaging material, so that colored images, free from black body absorbers, can be produced.

[0016] Accordingly, this invention provides a method for transferring a contrasting pattern of intelligence from an ablation-transfer imaging medium to a receptor element in contiguous registration therewith, the medium comprising:

- (i) a support substrate;
- (ii) at least one intermediate dynamic release layer essentially coextensive with the substrate; and
- (iii) a radiation-ablative topcoat, also essentially coextensive with the substrate,

the dynamic release layer and/or the topcoat containing at least one radiation-ablative binder, the dynamic release layer being capable of absorbing radiation at a rate sufficient to effect the imagewise ablation mass transfer of the topcoat to the receptor element, which method comprises imagewise irradiating the medium according to such pattern of intelligence with an intensity sufficient to effect the imagewise ablation mass transfer of the volume of the imagewise-exposed area of at least the topcoat of the medium securedly on to the receptor element. The method of this invention is characterized in that the dynamic release layer comprises a non-black body radiation absorber but is free from black body radiation absorber, and in that the topcoat comprises an imaging amount of a non-black body contrast imaging material, this contrast imaging material being transferred to the receptor element to delineate the pattern of intelligence thereon.

[0017] This invention also provides an ablation-transfer imaging medium comprising:

- (i) a support substrate;
- (ii) at least one intermediate dynamic release layer essentially coextensive with the substrate; and
- (iii) a radiation-ablative topcoat, also essentially coextensive with the substrate,

the dynamic release layer and/or the topcoat containing at least one radiation-ablative binder, the dynamic release layer being capable of absorbing radiation at a rate sufficient to effect the imagewise ablation mass transfer of the topcoat to the receptor element. The medium of this invention is characterized in that the dynamic

release layer comprises a non-black body radiation absorber but is free from black body radiation absorber, and in that the topcoat comprises an imaging amount of a non-black body sensitizing contrast imaging material.

FIGURE 1A is a side view photomicrograph of a composite imaging medium according to the present invention and the illuminated space above same, prior to laser exposure;

FIGURE 1B is a like photomicrograph taken under identical conditions, but 100 nanoseconds after initiation of a 260 nanosecond laser pulse directed through the support substrate of said composite imaging medium and into the dynamic release layer and ablative topcoat thereof;

FIGURE 2 is a schematic/diagrammatic illustration of the method/system according to the present invention, including one embodiment of the composite imaging medium wherein the support substrate thereof is transparent to the imaging radiation;

FIGURE 3 is a schematic/diagrammatic illustration of another method/system of this invention, including a second embodiment of the composite imaging medium wherein the support substrate thereof is not transparent to the imaging radiation; and

FIGURE 4 is a graph plotting transferred image density versus laser power and illustrates the greater imaging sensitivity attained using a composite imaging medium of the invention.

[0018] As already mentioned, the present ablation-transfer imaging medium necessarily includes at least one dynamic release layer (ii) intermediate the support substrate (i) and the imaging radiation-ablative carrier topcoat (iii).

[0019] By "dynamic release layer" is intended an intermediate layer that must interact with the imaging radiation to effect imagewise ablative transfer of at least the carrier topcoat onto a receptor element at an energy/fluence less than would be required in the absence thereof. The dynamic release layer is believed to release the carrier topcoat by effectively eliminating the adhesive forces that bond or consolidate the carrier topcoat with the support substrate. Preferably, under the same conditions additional propulsion is simultaneously provided by the interaction of the imaging radiation therewith, e.g., by ablation of the dynamic release layer itself, thus further facilitating the imagewise ablative transfer of the entire carrier topcoat to a receptor element.

[0020] The dynamic release layer (DRL) is at least one layer of any organic or inorganic material, or combination thereof, that absorbs at least a fraction of the imaging radiation sufficient to diminish the adhesion (in the imagewise exposed areas) between said at least one DRL and the support substrate, the DRL(s) and the carrier topcoat, or both. Such material can be intrinsically absorbing, or sensitized to absorb wavelengths of the imaging radiation. Preferably, the DRL(s) are intrinsically or inherently absorbing of the imaging radiation, highly absorbing of the imaging radiation such that very thin layers thereof can be employed, e.g., at least one low melting thin metal film. Further, these materials respond effectively to the imaging radiation on the nanosecond time scale, or even faster.

[0021] Exemplary such absorbing materials suitable for the DRL include thin films of metals, metal oxides, and metal sulfides which effectively melt, vaporize or otherwise change physical state when exposed to imaging radiation and preferably have little or no toxicity, have low energy requirements for release and reflect as little of the imaging radiation as possible. Representative such metals are those metallic elements of Groups Ib, IIb, IIIa, IVa, IVb, Va, Vb, Via, Vlb, VIb and VIII of the Periodic Table, as well as alloys thereof or alloys thereof with elements of Groups Ia, IIa, and IIIb, or mixtures of same. Particularly preferred metals include Al, Bi, Sn, In or Zn, and alloys thereof or alloys thereof with elements of Groups Ia, IIa and IIIb of the Periodic Table, or their mixtures. Suitable such metal oxides and sulfides are those of Al, Bi, Sn, In, 2n, Ti, Cr, Mo, W, Co, Ir, Ni, Pd, Pt, Cu, Ag, Au, Zr or Te, or mixtures thereof.

[0022] Other exemplary DRL materials include those that undergo a change in crystal structure upon exposure to imaging radiation, such as germanium or sensitized liquid crystal materials. Also exemplary are those materials described in US-A-4 756 633 and US-A-4 897 310.

[0023] Exemplary organic DRL materials comprise sublimable materials and monomeric and polymeric compounds which are intrinsically capable of absorbing imaging radiation, and/or monomeric and polymeric compounds which have been sensitized with non-black body absorbers to impart the necessary absorbance. Representative monomeric compounds include metal phthalocyanines, metal dithiolenes, anthraquinones, etc., which may be, e.g., vacuum deposited in the form of a thin layer. Representative polymeric compounds include the polythiophenes, polyanilines, polyacetylenes, polyphenylenes, polyphenylene sulfides, polypyrroles, and derivatives or mixtures thereof.

[0024] Yet other exemplary DRL materials include the combination of any ablation sensitizer/absorber, e.g., any near infrared or visible light sensitizer, in at least one binder which need not be, but may be absorbing at the imaging wavelength, or a sensitized/absorbing binder alone. Advantageously, such binder material is polymeric and preferably comprises those ablative polymers which undergo rapid acid catalyzed partial decomposition.

[0025] Depending upon the particular dynamic release material(s) selected and the imaging conditions employed, at least a portion of the DRL may itself be transferred together with the carrier topcoat onto the receptor element during the ablation process. However, since it contains a non-black body absorber, the dynamic release layer itself transfers essentially no contrasting material to the final product. This is particularly important when using the ablation-transfer

imaging media of the present invention for applications where color fidelity is important, e.g., for color printing and proofing.

[0026] The thickness of the at least one dynamic release layer depends upon the material(s) selected therefor. For example, where a metal such as those discussed above is employed as the DRL, a thickness of about one monolayer of the metal to 50nm (500 angstroms) is preferred.

[0027] The dynamic release layer preferably synergistically interacts with the ablative carrier topcoat to reduce the threshold energy required for transfer. This enables complete carrier topcoat transfer at lower energy inputs thereby requiring only small amounts, and even none at all, of an ablation sensitizer in the carrier topcoat as more fully described below, and also enables the more effective utilization of any other functional additives, also described below.

[0028] The composite ablation-transfer imaging media of the present invention also necessarily comprise an imaging radiation-ablative carrier topcoat (iii) essentially coextensive with the support substrate (i) and the at least one DRL (ii).

[0029] Such ablative carrier topcoat itself necessarily contains an imaging amount of a contrast imaging material.

[0030] By "contrast imaging material" is intended that material used to distinguish the resulting pattern of intelligence transferred to the receptor element.

[0031] Such contrast imaging material, moreover, either may or may not itself be an ablation sensitizer capable of promoting ablation-transfer under the intended imaging conditions that result in ablation. Failure of the contrast imaging material to itself initiate or promote ablation (i.e., a "non-ablation sensitizing contrast imaging material") may be the result of a lack of absorbance at the ablation wavelength(s), a lack of sufficient absorbance of same, or a failure of absorbance to result in a pressure build up phenomenon, e.g., the absorbance provides a non-ablation promoting event such as photobleaching, stable triplet, fluorescence or phosphorescence. Thus, the contrast imaging material must be visible or discernible to the detector/technique used to distinguish the resulting pattern of intelligence transferred to the receptor element and/or remaining on the imaging medium, per se.

[0032] Exemplary such contrast imaging materials that can be ablatively transferred to a receptor element in a pre-determined contrasting pattern of intelligence to visibly or symbolically represent or describe an object or data include colorants (dyes or pigments), ultraviolet and infrared absorbing materials, polymeric materials, magnetic materials, fluorescent materials, electrically conducting materials, etc.

[0033] Still other additives may be included to enhance the film properties and transfer characteristics. These additives need not function as a contrast imaging material and include, e.g., plasticizers, flow additives, slip agents, light stabilizers, anti-static agents, surfactants, brighteners, anti-oxidants and others known to the formulation art.

[0034] In a preferred embodiment of the present invention, the subject ablation-transfer imaging/recording technique is advantageously photo- and more preferably laser-induced, albeit it need not be (for example, any light source of sufficient intensity in the near infrared or visible spectral region is suitable).

[0035] Photo- or laser-induced ablation-transfer comprehends a threshold energy below which no effective material transfer occurs and a requirement that the energy be input at a rate greater than the ability of the materials to reverse the factors leading to the aforementioned pressure accumulation, for example by excessive thermal diffusion outside the irradiated area. Thus, imaging radiation capable of exceeding the threshold energy (fluence, joules/cm²) and power density (watts/cm²) is required for effective image transfer. By proper selection of materials and imaging parameters, this latter requirement can lead to exposure times on a nanosecond time scale which is at least ten times faster than exposure times necessary for conventional transfer imaging processes. The actual values of fluence and power density suitable for photo- and laser-induced ablative transfer imaging are dependent on the specific materials employed in the imaging medium and the specific receptor selected. In addition, fluence is power density dependent, particularly in the event that thin metal DRLs are employed.

[0036] In the present invention, the dynamic release layer and/or the topcoat contains at least one radiation-ablative binder (also referred to herein as an "ablative binder"). The ablative binder need not be absorbing at the imaging wavelength but is essentially completely imagewise mass transferred to the receptor element. Suitable binders include those materials, e.g., polymeric materials, which are effective in adhering the carrier topcoat to the DRL prior to transfer, as well as adhering the transferred material to the receptor element after exposure to imaging radiation.

[0037] Advantageously, the ablative binder material referred to above comprises a conventional film-forming polymer providing high visible transparency. Exemplary such film-forming polymers include, but are not limited to, polycarbonates, polysulfones, styrene/acrylonitrile polymer, polystyrenes, cellulosic ethers and esters, polyacetals, polymethylmethacrylate, polyvinylidene chloride, α -chloroacrylonitrile, maleic acid resins and copolymers, etc.

[0038] In particular, such ablative binders according to this invention are those binders which decompose rapidly to produce effective amounts of gases and volatile fragments at temperatures of less than 200°C as measured under equilibrium conditions and the decomposition temperatures of which are significantly reduced in the presence of small amounts of generated acids. Most preferably, the decomposition temperatures thereof are decreased to less than about 100°C.

[0039] Exemplary such polymers include nitrocellulose, polycarbonates and other polymers of the type described in J.M.J. Frechet, F. Bouchard, F.M. Houlihan, B. Kryczke and E. Eichler, J. Imaging Science, 30(2), pp. 59-64 (1986),

and such other polymers as polyurethanes, polyesters, polyorthoesters and polyacetals, and copolymers thereof.

[0040] In a preferred embodiment of the invention, the carrier topcoat also includes, in addition to the ablative binder, an ablation sensitizer (absorber). By "ablation sensitizer" is intended any initiator capable of initiating and promoting the ablation process. It does this by absorbing the imaging radiation and transferring the absorbed energy into an explosive ablative force. Such types of sensitizers/ initiators are well known to the recording art as discussed earlier in the context of optical disk writing. Light sensitization for imaging materials is of course also well known to the recording art.

[0041] In another preferred embodiment of the invention, the imaging radiation-ablative carrier topcoat comprises at least one ablation sensitizer which absorbs at the wavelength of the desired laser output, for example in the near infrared spectral region of 760 nm to 3,000 nm, and at least one ablative binder, the at least one sensitizer being present in an amount sufficient to effect the rapid partial decomposition of the at least one binder when the at least one sensitizer interacts with laser light. The ablative binder advantageously comprises those polymeric materials which undergo rapid acid catalyzed partial decomposition, preferably at temperatures less than 200°C as measured under equilibrium conditions. The carrier topcoat may also, optionally, contain materials which are non-absorbing at the wavelength of the desired laser output and/or non-decomposing.

[0042] In another embodiment of the invention, the ablative binders are advantageously those polymeric materials which transfer under the imaging conditions, and are preferably (especially when the DRL comprises an organic absorber), those which undergo rapid acid catalyzed partial decomposition at temperatures of less than 200°C as measured under equilibrium conditions, and most preferably at temperatures of less than about 100°C as measured under equilibrium conditions.

[0043] In another embodiment of the invention, the carrier topcoat comprises at least one near infrared sensitizer, at least one ablative binder, and at least one hydrogen atom donating material (H•) for the acid catalyzed decomposition of the ablative binder (which may be present in the binder itself). These latter materials include alcohols, thiols, phenols, amines and hydrocarbons.

[0044] In another preferred embodiment of the present invention, a near infrared laser-ablation transfer imaging medium is provided. Such medium advantageously comprises a near infrared transparent support film bearing at least one DRL and a carrier topcoat of near infrared ablative coating containing a substantially colorless near infrared sensitizer. This medium can be effectively and advantageously employed for color imaging when a colorant is added.

[0045] Upon exposure to laser light, the DRL and the absorbing sensitizer interact with laser light to cause rapid release of at least the carrier topcoat and may involve partial decomposition of the DRL and/or carrier topcoat binders to gaseous and non-gaseous products. The rapid expansion of the heated gases assists ablation of the exposed carrier topcoat onto an adjacent receptor sheet providing a reverse of the imaged color film (i.e., a color print or proof).

[0046] Suitable absorbing sensitizers according to the present invention include any material which can absorb at a desired wavelength for a particular near infrared or visible imaging wavelength. In particular, where visibly transparent coatings are required, for example, substituted aromatic diamine dication diradical type sensitizers or cation radical sensitizers with counterions derived from strong acids and absorbing in the near IR are preferred.

[0047] Examples of these sensitizers include the series of near infrared absorbers marketed under the trademarks Cyasorb IR 165, 126 and 99 by American Cyanamid, as well as those IR absorbers described in US-A-4 656 121.

[0048] Radiation sources emitting near infrared wavelengths in combination with visibly colorless sensitizers are preferred for high fidelity color imaging applications. In other applications, any radiation source of sufficient intensity, typically not less than 10⁴ watts/cm², emitting in the visible and/or near infrared can be employed for photo-ablation without limitation to black body sensitizers as essentially required by the prior art. The sensitizers of the present invention are most preferably highly absorbing at the wavelengths of the imaging radiation and soluble in the binders employed. Examples of suitable non-black body sensitizers which can be effectively employed in the ablative topcoat are cyanine dyes, phthalocyanine dyes, metal dithiolenes, methylene blue salts, di- and triarylmethane cation salts, Wurster's blue salts, and other visibly or near infrared absorbing onium salts derived from strong acids, etc. Various of these are described in US-A-4 315 983, US-A-4 508 811, US-A-4 948 776, US-A-4 948 777, US-A-4 948 778 and US-A-4 950 640.

[0049] The support substrates (i) employed can be either support films transparent to the imaging radiation or non-transparent such support films. Transparent support films which can be employed include glass, polyesters (by reason of their high optical clarity and dimensional stability), polycarbonates, polyurethanes, polyolefins, polyamides, polysulfones, polystyrenes, cellulose and any support substrate which does not dissolve in the coating solvents employed, with polyesters being preferred. Examples of non-transparent supports include any non-transparent support substrate which would not dissolve in the coating solvents employed. These supports can include filled and/or coated opaque polyesters, aluminum supports, such as used in printing plates, and silicon chips. The thickness of such support substrates is not critical and can vary widely, same depending, for example, upon the particular intended application and whether irradiated from the front or back surface thereof.

[0050] An anti-reflection layer (vis-a-vis the imaging radiation) may optionally be provided on the face surface of the

support opposite the ablative carrier topcoat and/or on the receptor element, to enhance efficiency of the ablative transfer by enabling more of the imaging radiation to be effectively utilized.

[0051] Such anti-reflection layer advantageously comprises one or more materials which are recognized for this purpose in the recording art, for example those described in US-A-3 793 022 and US-A-4 769 306, and are also applied in known manner. Suitable materials, e.g., for a polyester support substrate having a refractive index of about 1.6, are coatable materials having refractive indices of about 1.3 to 1.4. Exemplary such materials include Fluorad (Registered Trademark) FC-721 from 3M Co., CaF_2 , MgF_2 , fluoropolymer, etc. The thickness of the anti-reflection layer(s) is selected as to provide the desired refractive properties for the layer(s) with respect to the wavelengths of the imaging radiation. For example, where Fluorad FC-721 is employed as the anti-reflection layer and 1064 nm imaging radiation is used, a thickness of 0.2 to 0.25 microns is effective.

[0052] To fabricate the ablation-transfer imaging media of this invention, the dynamic release layer is deposited onto appropriate support substrate by any means well known to this art, e.g., vapor deposition, sputtering, wet deposition, etc.

[0053] The DRL side of the resulting intermediate article is then overcoated with a solution or dispersion of the constituents of the carrier topcoat by coating techniques also well known to this art, such as Meyer rod coating, gravure coating, reverse roll coating, modified bead coating, extrusion coating, etc., and the coated article is dried. The coating solution or dispersion contains solvent(s), the contrast imaging material, any binder(s) and sensitizer(s), additives, etc., depending upon the intended final application.

[0054] The solvents employed in the present invention are preferably those which dissolve both the binders and the optional sensitizers. Exemplary such solvents include, but are not limited to, water, alcohols, esters, ketones, aliphatic and aromatic hydrocarbons, chlorinated aliphatic and aromatic hydrocarbons, nitro aliphatic and aromatic hydrocarbons, amides and others known to the formulation art.

[0055] The thickness of the dynamic release layer(s) is optimized to provide for the total transfer of the carrier topcoat to the receptor element with a minimum energy requirement (joules/cm²). The thickness of the carrier layer is optimized to provide the amounts of material required for a particular application; for example, an amount of dye or pigment necessary to meet a density requirement on the receptor element or an amount of UV absorber required to meet a UV blocking specification of the donor medium or mask.

[0056] The composite imaging media of the present invention are useful for a wide variety of applications, including color printing and color proofing. Other than these, and depending upon the nature of the contrast imaging material contained in the ablative carrier topcoat (whether or not also containing an ablation sensitizer), the subject composite imaging media may include a UV absorber and thus be adopted for the production of exposure masks, and infrared dye for generating machine readable images, e.g., bar codes, magnetic particles for preparing machine readable documents, fluorescent materials for coding security documents, or electrically conductive materials for the production of electronic components, e.g., printed circuit boards and other direct-to-plate and direct-to-board applications, etc.

[0057] In one embodiment of this invention, providing an imaging medium well suited, e.g., for color proofing, the carrier topcoat includes a sensitizer selected as to be substantially non-absorbing in the visible spectral region (380-760 nm) and a pigment dispersion of a colorant supplied as a nitrocellulose based printing ink. The sensitizer is present in an amount of about 0.05% to about 50% by weight of dry solids. Preferably, the sensitizer is present in an amount of 2% to 25% by weight of dry solids. The total amount of dry solids (grams/meter²) is adjusted to meet American Newspaper Publishers Association (A.N.P.A.) or Specifications for Web Offset Publications (S.W.O.P.) density specifications for solid color densities transferred to the receptor element, e.g., known commercial newsprint. The dynamic release layer is advantageously a thin metal film applied to a support substrate, e.g., a polyester film. Preferably, the thin metal film is a low melting, low-toxicity metal or metal alloy of Al, Bi, Sn, In and/or Zn. The thickness of the metal film advantageously ranges from one monolayer to 50 nm (500 Angstroms), the optimal thickness depending on the particular metal. By "color proofing" is of course intended that technique, very well known to the recording art, of predicting or confirming one or more aspects of color printing prior to press, e.g., color rendition, tonal rendition, registration, composition, and the like.

[0058] In another embodiment of the composite imaging medium of the invention, also well suited for color proofing and printing applications, the carrier topcoat is also deposited onto the DRL (and then dried) as a pigment-dispersion of a colorant, i.e., a nitrocellulose based printing ink, but contains no sensitizer. The dynamic release matrix is again a thin metal film applied onto a polyester film support substrate. Preferably, the thin metal film is a low melting, low-toxicity metal or metal alloy of Al, Bi, Sn, In and/or Zn. The thickness of the metal film can again range from one monolayer to 50 nm (500 Angstroms), the optimal thickness depending on the particular metal employed. In this embodiment, the receptor element to which the proof/print is transferred, by reason of the absence of the ablation-promoting sensitizer, is advantageously passively adhesively coated, e.g., with a hot melt adhesive, to assist securing the ablated image thereto.

[0059] In another embodiment, i.e., masking, in which the imaging media can be effectively employed as an exposure mask for use in graphic arts or printed circuit preproduction, the contrast imaging material is an absorber which is

effective in blocking the light output from common exposure devices. Exemplary such materials are azo derivatives, oxediazole derivatives, dicinnamalacetone derivatives, benzophenone derivatives, etc. By "masking" is intended that operation, also very well known to the recording art, including blanket exposure of a typically light sensitive material, e.g., printing plate, resist, diazo, etc., through a pre-existing pattern of intelligence, e.g., a "mask", which selectively blocks the exposure radiation according to the pattern of intelligence, e.g., a printed circuit, newspaper page, etc. One specific example of an imaging medium well suited for such applications comprises a polyester film support substrate, a thin metal film dynamic release layer, e.g., an Al, Bi, Sn, In and/or Zn thin film, or thin film of an alloy thereof, in a binder polymer, and an ablative carrier topcoat comprising a nitrocellulose based printing ink vehicle, curcumin, and, optionally, a sensitizer, e.g., a cation radical or a dication diradical derivative of a triarylamine compound.

[0060] In still another embodiment, in which the imaging media can be effectively employed in a security printing application, the contrast imaging materials are substantially colorless, but which fluoresce in the visible region when exposed to ultraviolet light. Representative such materials include oxazole derivatives, oxediazole derivatives, coumarin derivatives, carbostyryl derivatives, etc. One specific example of an imaging medium well suited for such applications comprises a polyester film support substrate, a thin metal film dynamic release layer, e.g., an Al, Bi, Sn, In and/or Zn thin film, or thin film of an alloy thereof, and an ablative carrier topcoat comprising a nitrocellulose based printing ink vehicle, bis(4-diethylaminophenyl)oxadiazole, and, optionally, a sensitizer, e.g., a cation radical or a dication diradical of a triarylamine compound.

[0061] In another embodiment, the contrast imaging material is readable by means of a near infrared scanner, e.g., a bar code. Exemplary such materials for this particular embodiment include metal dithiolene derivatives, phthalocyanine derivatives, triarylamine derivatives, graphite, carbon black, chromium dioxide and mixtures thereof. One specific example of an imaging medium well suited for these applications comprises a polyester film support substrate, a thin metal film dynamic release layer, e.g., an Al, Bi, Sn, In and/or Zn thin film, or thin film of an alloy thereof, and an ablative carrier topcoat comprising a nitrocellulose based ink vehicle, a phthalocyanine derivative, and, optionally, a sensitizer, e.g., a cation radical or a dication diradical of a triarylamine compound.

[0062] In yet another embodiment, the contrast imaging material is magnetic, for the production of such machine readable items as information strips, checks, credit cards, etc. Exemplary thereof are iron, iron oxide, cobalt-iron oxide, barium ferrite, mixtures of the above, and the like.

[0063] In still another embodiment, the contrast imaging material includes at least one additive which comprises a hydrophobic ink accepting resin for transfer to a grained aluminum surface used, for example, in the production of offset lithographic plates. Suitable additives for this application comprise phenolformaldehyde resins, crosslinked polyvinyl alcohol resins, crosslinked epoxy resins, etc.

[0064] In a further embodiment, the contrast imaging material includes at least one additive which comprises an electrically conductive material for transfer to an insulating surface used, for example, in the production of printed circuits. Suitable additives for this application comprise conductive dispersions of silver, aluminum, copper, etc.

[0065] In each of the above embodiments, the imaging medium preferably includes an anti-reflection layer, as above described, on the face surface of the support substrate opposite the DRL and comprising, for example, a fluoropolymer, magnesium fluoride or calcium fluoride.

[0066] The material transfer phenomenon of the laser-induced ablation process of the present invention is illustrated by the photomicrographs obtained by time resolved grazing incident microscopy, TRGIM, in Figures 1A and 1B.

[0067] FIGURE 1A is a side view photomicrograph of a composite imaging medium of the invention and the illuminated space thereabove (in lieu of a receptor element), prior to laser exposure.

[0068] FIGURE 1B is a like photomicrograph, taken 100 nanoseconds after the initiation of a 260 nanosecond pulse from a Nd:YAG laser, 5, directed through the polyester support, 4, about 6 mm (¼ inch) from the edge thereof and into the dynamic release layer, 3, and absorbing ablative topcoat, 2 (the imaging medium, per se, is more fully described in the Example 1 which follows), to produce a plume, 6, of ablated materials. The horizontal lines, 1, in the space above the medium are interference lines resulting from the use of coherent probe illumination. The Nd:YAG laser (1064 nm output from a Quantronics 116FL-O laser controlled by an Anderson Laboratories' DLM-40-IR2.7 acousto-optic modulator and a 40 MHz signal processor) delivered 0.6 J/cm² in a 25 micron diameter beam (1/e²).

[0069] FIGURE 2 illustrates the use of an imaging radiation transparent support substrate in the method of the present invention. In this embodiment, imaging radiation, 6, impinges upon the imaging material which comprises an anti-reflection first layer, 5, an imaging radiation transparent support substrate, 1, a dynamic release layer, 2, and an ablative carrier topcoat, 3. Also shown is a receptor element, 4, in contiguous registration therewith.

[0070] FIGURE 3 illustrates an alternative embodiment of the present invention wherein the imaging material comprises a nontransparent support substrate, 7. In this embodiment, the receptor element, 4, is made of imaging radiation transparent material and is provided with an anti-reflection back layer, 5. The imaging radiation, 6, impinges upon the imaging material from the front or through the receptor element, into the ablative carrier topcoat, 3, and dynamic release layer, 2.

[0071] In either embodiment, a pattern of imaging radiation at the desired wavelength(s) is directed into the absorbing

layer(s). This causes ablation of at least the carrier topcoat and its transfer to the receptor element, thus producing an imaged donor film, 8, and a corresponding image of opposite sign, 9, on the receptor element.

[0072] The imaging radiation employed in this invention includes wavelengths in the visible and near infrared spectral regions. Near infrared radiation is preferred for color proofing and printing applications because the sensitizers, when indeed they are employed for a particular application, can be substantially colorless in the visible region and thus will not adversely affect the color fidelity of the transferred image on the receptor element.

[0073] Exemplary radiation emitting devices for imagewise exposing the composite imaging media of the invention include solid state lasers, semiconductor diode lasers, gas lasers, dye lasers, xenon lamps, mercury arc lamps, and other visible and near infrared radiation sources which are capable of providing sufficient energy to equal, or exceed, the threshold energy for ablation transfer and of providing this energy at such a rate as to institute that phenomenon of transient pressure accumulation discussed earlier and believed responsible for the ablative transfer process.

[0074] The actual value of threshold energy is intensity dependent as well as materials dependent. Typically, when employing imaging media constructions comprising a dynamic release layer composed of high thermal conductivity thin metal films, such as aluminum, at least 50 mJ/cm² at 10⁶ watts/cm² is required, while at least 100 mJ/cm² at 10⁴ watts/cm² is more typical for organic constructions. When these conditions are not satisfied, undesirable processes rather than, or in addition to, ablation-transfer may occur, e.g., melting, sublimation, charring, etc. Incomplete transfer and/or image quality degradation may result. Other constraints on the exposure device include the ability to focus the imaging radiation to a desirable spot size and depth and to modulate the radiation at dwell times suitable for the desired imaging application.

[0075] Particularly representative devices for providing the imaging radiation include lasers such as Nd:YAG lasers emitting at 1064 nm, for example that incorporated in the imaging hardware of the Crosfield Datrix 765 laser facsimile writer, laser diode systems emitting at 780-840 nm, or other radiation sources designed to provide a power density of 10⁴ watts/cm² or greater.

[0076] The radiation source is preferably focused to provide the most efficient utilization of energy when it is impinged upon the imaging medium.

[0077] The receptor element need not be specially treated or selected to effectively receive materials from the donor medium and can include, for example, those which are well-known in the art of proofing and printing, such as newsprint, coated or uncoated papers of all shades and color, opaque filled and opaque coated plastic sheets, with the printing stock to be employed in the particular color proofing application being preferred. Other suitable receptors include fabrics, wood, cardboard, glass, ceramics, leather, metals such as aluminum and copper, rubber, papers and plastics generally, etc. While the receptor element need not be specially treated or selected to assist in the ablation-transfer process, as indicated above, it is nonetheless also within the scope of this invention to employ a treated or coated receptor, for example a receptor sheet coated with an effective amount of an adhesive or sizing, to aid in the adhesion of the ablated carrier topcoat thereto at lower energy than otherwise would be required.

[0078] The imaging medium is most advantageously positioned and firmly maintained in face-to-face registered direct contact with the particular receptor element selected, to facilitate transfer ablated carrier topcoat thereto, by any means suitable for such purpose, e.g., positive pressure, a vacuum, or even by the adhesive properties of the receptor element itself.

[0079] From the foregoing, it will be seen that the present invention provides numerous advantages and a versatility hitherto alien to this art. Primarily, by reason of the DRL interlayer, the subject composite imaging media are significantly more sensitive to imaging radiation, certain constructions being at least twice as sensitive using far less sensitizer, e.g., at least 75% less, at about half the thickness of the ablative topcoat, to achieve a comparable image density on the receptor element. Correspondingly, the energy/power requirements for a clean transfer are significantly decreased. Moreover, the ablative carrier topcoat need not itself be sensitized, albeit this is optional; rather, only the dynamic release layer is required to be sensitive to the imaging radiation.

[0080] In like vein, selection of binder material is expanded and any required amount thereof lessened per the present invention, as decomposable binders are not required, as is the selection of appropriate sensitizers.

[0081] Further, the distribution of energy absorption in the DRL and ablative topcoat is optimized per the present invention and the ablative process is kinetically much faster, a not insubstantial attribute, considering that this permits much faster access to the desired final product.

[0082] It too will be appreciated that a very primary advantage of the invention is its flexibility in enabling virtually any type of topcoat material/pattern of intelligence to be transferred to virtually any receptor element.

[0083] In order to further illustrate the present invention and the advantages associated therewith, the following specific examples are given, it being understood that the same are intended only as illustrative and in nowise limitative.

[0084] In said examples to follow, all parts and percentages are given by weight, unless otherwise indicated.

EXAMPLE 1:

[0085] This example describes the production of various composite ablation-transfer imaging media of the present invention and also the use of same to prepare a newspaper multi-color proof.

[0086] Near infrared sensitized formulations of cyan, magenta and yellow gravure (flexo) ink dispersions (marketed by American Inks & Coatings Corp.), as well as a black, graphite colloidal suspension (custom made by Acheson Colloids Co. and described in US-A-4 245 003) were separately machine coated by a modified bead coating technique, to a dry thickness of about 0.5 μm , the three colorant inks onto the metallized face surfaces of three 644 mm (25 inch) wide webs of 50% transmission aluminized 117 μm (4.6 mil) ICI Melinex 505 polyester film to produce three imaging media and the black suspension onto a 644 mm (25 inch) web of 117 μm (4.6 mil) ICI Melinex 516 polyester film to produce a fourth imaging medium.

Carrier Topcoat Coating Formulations:

[0087] Near Infrared Sensitizer (1064 nm): Cyasorb (Registered Trademark) IR 165 (American Cyanamid Corp., Glendale Protective Technologies Division).

Cyan Dispersion:

[0088]

2.44 parts by weight Ink (93-405 marketed by American Inks & Coatings),
0.08 parts by weight IR 165,
19.4 parts by weight Solvents (50% Methyl Ethyl Ketone (MEK), 20% Methyl Propyl Ketone (MPK), 15% n-Butyl acetate (n-BuAc) & 15% Cyclohexanone).

Magenta Dispersion:

[0089]

2.88 parts by weight Ink (93-418 marketed by American Inks & Coatings),
0.08 parts by weight IR 165,
23.2 parts by weight Solvents (50% MEK, 20% MPK, 15% n-BuAc & 15% Cyclohexanone).

Yellow Dispersion:

[0090]

2.62 parts by weight Ink (93-4149 marketed by American Inks & Coatings),
0.08 parts by weight IR 165
21.9 parts by weight Solvents (50% MEK, 20% MPK, 15% n-BuAc & 15% Cyclohexanone).

Black Suspension: (not part of present invention)

[0091]

7.5 parts by weight Graphite Suspension obtained from Acheson Colloids Co.,
20.5 parts by weight. Solvents (50% MEK, 20% MPK, 15% n-BuAc & 15% cyclohexanone).

[0092] The dry optical transmission densities of the imaging media thus produced, determined using a Macbeth Densitometer (TD-904), were: CYAN = 0.72 density units (d.u.); MAGENTA = 0.63 d.u.; YELLOW = 0.73 d.u.; BLACK = 1.39 d.u.

[0093] A four color proof was produced on a Crosfield Datrax 765 reader/writer from the Color Test Negative color separations provided by the American Newspaper Publishers Association (A.N.P.A.). The respective color-separated images were scanned by the reader and converted to registered digital electronic signals and transmitted sequentially to the laser writer, a scanning YAG laser adjusted to write 47.3 lines mm^{-1} (1,200 lines per inch) at 7.5 watts (fluence 160 mJ/cm^2 , power density 10⁶ watts/ cm^2 , beam diameter 25 microns (1/e²), dwell time about 104 nanoseconds). A proof or print was produced by serially and successively placing the aforesaid four imaging media in direct face-to-

face contact with a common newspaper stock receptor sheet securely maintained in place by vacuum in the laser writer and respectively imaging/exposing same.

[0094] The carrier topcoats of the four imaging media were imagewise ablated to the newspaper receptor sheet in primary color areas at the following optical reflection densities (measured using a X-Rite 428 densitometer):

CYAN: 0.96 d.u. vs. 0.90 d.u. A.N.P.A. specification
MAGENTA: 0.89 d.u. vs. 0.90 d.u. A.N.P.A. specification
YELLOW: 0.87 d.u. vs. 0.85 d.u. A.N.P.A. specification
BLACK: 1.12 d.u. vs. 1.05 d.u. A.N.P.A. specification

[0095] The 406 x 609 mm (16 x 24 inch) multi-color proof thus produced, in about 10 minutes, corresponded favorably to a newspaper press print produced from printing inks containing the same pigments and printing plates produced from the same color-separation negatives.

EXAMPLE 2:

[0096] This example describes the production of various composite ablation-transfer imaging media of the present invention and also the use of same to prepare a magazine multi-color proof.

[0097] The dispersions/suspension prepared as in Example 1 were coated onto the metallized face surfaces of aluminized polyester films of 50% transmission by hand drawdown using a #7 Meyer rod (cyan), a #5 Meyer rod (magenta) and #3 Meyer rod (yellow). The black suspension was hand coated onto ICI Melinex 516 polyester film using a #9 Meyer rod. The four imaging media were sequentially ablated (Datrax writer adjusted to 47.3 lines mm⁻¹ (1,200 lines per inch) resolution and 8.5 watts, fluence 180 mJ/cm²) onto a #5 ground wood receptor sheet (Ad-Proof Enamel (Trade Mark) from Appleton Paper Inc.) of about 70% brightness and about 27.2 Kg (60 lb.) basis weight, by the technique of Example 1, to provide the following optical reflection densities:

CYAN: 1.50 d.u. vs. 1.50 d.u. A.N.P.A. specification
MAGENTA: 1.43 d.u. vs. 1.45 d.u. A.N.P.A. specification
YELLOW: 1.05 d.u. vs. 1.05 d.u. A.N.P.A. specification
BLACK: 1.30 d.u. vs. 1.60 d.u. A.N.P.A. specification

EXAMPLE 3:

[0098] This example relates to providing an anti-reflection (AR) layer, or coating, on the back surface of a composite ablation-transfer imaging medium of the invention and reports certain comparative data obtained by ablating the AR coated and uncoated imaging media onto two different substrates.

[0099] The uncoated, raw polyester side of a machine coated, 203 x 254 mm (8 x 10 inch) magenta composite imaging medium prepared as in Example 1 was coated with Fluorad FC-721, a 3M fluorochemical coating solution having a refractive index of about 1.36, to a thickness of about 0.2 µm to 0.25 µm (cross-sections were measured with a scanning electron microscope) using a #4 Meyer rod. The AR coated imaging medium and an uncoated control imaging medium of the same sample were ablatively imaged onto plain, untreated white paper and onto newspaper stock using a Crosfield Datrax 765 Nd:YAG laser writer. The comparative results obtained are reported in Table I below:

TABLE I:

IMAGING MEDIUM	WATTS AT 47.3 lines mm ⁻¹ (1200 lines/inch)	RECEPTOR	% MAGENTA INK COATING TRANSFERRED
AR Coated	4.0	plain paper	38.5

TABLE I: (continued)

IMAGING MEDIUM	WATTS AT 47.3 lines mm ⁻¹ (1200 lines/inch)	RECEPTOR	% MAGENTA INK COATING TRANSFERRED
Uncoated	4.0	plain paper	10
AR Coated	5.0	plain paper	100
Uncoated	5.0	plain paper	41.4
AR Coated	5.0	newsprint	100
Uncoated	5.0	newsprint	50.5
Uncoated	5.5	newsprint	100

EXAMPLE 4:

[0100] This example compares the imaging properties of a variety of binder polymers constituting the ablative carrier topcoat of the composite ablation-transfer imaging media according to the invention.

[0101] The imaging properties of a variety of solvent soluble polymers were compared. Each resin (0.2 grams) was added to 9.8 grams of a mixture containing 0.3% of 2,5-dimethyl-3-hexyne-2,5-diol, 0.7% Cyasorb IR 165, and 5% of Morfast Yellow 101 by weight. The solvents selected were a 50:50 mixture of methylene chloride and 1,1,1-trichloroethane or a 50:20:15:15 mixture of methyl ethyl ketone, methyl propyl ketone, n-butyl acetate and cyclohexanone. Appropriate corrections were made for the density differences between the two solvent blends.

[0102] Films were prepared by coating the above solutions onto 50% transmission aluminized polyester with a #3 Meyer rod, to a transmission density (blue filter) of .60. The films were then ablated at three power settings on a Crosfield Datrax 765 Writer with a Nd:YAG laser onto an uncoated white paper to provide a maximum reflectance density of 0.75.

[0103] The following Table II reports the results of this experiment as the amount transferred at three power settings expressed as a percentage calculated by:

$$\text{measured density}/\text{maximum density} \times 100\%.$$

TABLE II:

Binder Polymer	Power (Watts) Fluence (Joules/cm ²)		
	6W 0.13 J/cm ²	4W 0.09 J/cm ²	3W 0.06 J/cm ²
Frechet Polycarbonate	100%	75%	45%
Bisphenol A Polycarbonate	100%	75%	0%
Polysulfone	100%	50%	0%
Styrene/Acrylonitrile	100%	65%	0%
α -Methylstyrene	100%	71%	25%
0.5 sec Cellulose Propionate	93%	16%	0%
Ethylcellulose	94%	36%	0%
Polystyrene	100%	64%	20%
Butvar TM	87%	25%	0%
Polymethylmethacrylate	96%	33%	0%
Unirez TM 7093	100%	70%	20%
Saran TM 120	94%	48%	0%
α -Chloroacrylonitrile	90%	30%	0%
Nitrocellulose SS	100%	75%	30%

EXAMPLE 5:

[0104] Imaging sensitivity plots (Laser Power vs. Cyan Density Transferred to the Receptor Sheet) derived from the cyan imaging medium and receptor sheets A as described in Example 1, and a similar cyan imaging medium/receptor B organization, but the latter imaging medium being devoid of a dynamic release layer, were compared. A Datrax writer

was adjusted to print solid patches of cyan colorant, i.e., maximum density, at 47.3 lines mm⁻¹ (1,200 lines/inch) for 3 watts, 4 watts, 5 watts, 6, watts, 7 watts and 8 watts of power. Average cyan densities, plus paper, were measured on each receptor at each power setting and plotted as shown in the graph of FIGURE 4.

EXAMPLE 6:

[0105] This example describes the production of various composite ablation-transfer imaging media of the present invention, the carrier topcoat thereof being devoid of any sensitizer, and also the use of same to prepare a multi-color proof on an adhesively coated receptor element.

[0106] The cyan (C), magenta (M), and yellow (Y) carrier topcoating formulations of Example 1, but without any IR 165 sensitizer being added thereto, were coated by the method described onto the metallized face surface of 50% transmission aluminized 117 μ m (4.6 mil) ICI Melinex 505 polyester film. These imaging media plus the black (K) imaging medium of Example 1 were used to ablate a four color print onto a specially coated receptor sheet using the same imaging/exposure device and under the same writing conditions of Example 1. The receptor sheet was prepared by sequentially coating a plain white paper with 3M Spray Mount (Registered Trade Mark) adhesive prior to the successive ablative transfer of images from the respective imaging media, in the order Y,M,C,K. Finally, a protective clear polyester sheet was laminated to the finished print.

EXAMPLE 7:

[0107] Magenta formulations according to Example 1 were prepared, both with and without the addition of the IR 165. Both formulations were coated as in Example 1 to provide a transmission density of 0.60 ± 0.05 units (green filter). The following receptor sheets were used:

1. Kimdura synthetic paper (Kimberly-Clark) as received.
2. Kimdura synthetic paper overcoated with a hot melt adhesive. The overcoating composition included the hot melt adhesive, i.e., Elvax 40W (Dupont, ethylene/vinyl acetate copolymer), 1 gram, and the tackifier, Foral 105 (Hercules, modified rosin ester), 1 gram, as well as a solvent blend (toluene/isopropanol, 90:10), 8 grams, and was applied by hand drawdown using a #10 Meyer rod, and air dried.

[0108] Magenta imaging media with and without sensitizer were imagewise ablated as described in Example 1 using the magenta film separation from the color test negative supplied by the American Newspaper Publisher's Association. The results and reflection densities reported in the following Table III were obtained:

TABLE III:

Imaging Medium	Receptor 1	Receptor 2
Magenta film without IR 165	no adhesion to receptor	0.90 ± 0.05 at 100% dot
Magenta film with IR 165	0.90 ± 0.05 at 100% dot	1.00 ± 0.05 at 100% dot

EXAMPLE 8:

[0109] A mask was prepared by coating the following formulation:

- (i) 2.70 grams Hittamine Arctic White Sol. (T1413) (Mobay Chemical Corp.), an ultraviolet absorber;
- (ii) 6.00 grams topcoat lacquer (American Inks and Coatings Corp. product #93-483);
- (iii) 20.0 grams solvent (50% MEK, 20% MPK, 15% n-BuAc, 15% cyclohexanone); and
- (iv) 0.21 grams Cyasorb IR 165 (American Cyanamid), onto the metallized face surface of a 50% transmission aluminized polyester film as described in Example 1 using a #5 Meyer rod.

[0110] After air drying, the coated film was imagewise ablated onto a porous paper receptor sheet using a Datrix YAG laser writer operating at 8 watts with 47.3 lines mm⁻¹ (1,200 lines/inch) resolution. The resultant mask had a Dmin = 0.15 and a Dmax = 3.40 (18A filter).

EXAMPLE 9:

[0111] Another type of mask was prepared by coating the following formulation:

- (i) 4.00 grams topcoat lacquer (American Inks and Coatings Corp. product #93-483);
- (ii) 2.10 grams Curcumin (Aldrich, CAS #458-37-7), an ultraviolet absorber;
- (iii) 0.20 grams Cyasorb IR 165 (American Cyanamid); and
- (iv) 15.0 grams solvent (50% MEK, 20% MPK, 15% n-BuAc, 15% cyclohexanone),

onto a 50% transmission aluminized polyester film as described in Example 1 with a #5 Meyer rod.

[0112] After air drying, the coated film was imagewise ablated onto a porous paper receptor sheet using a Datrax 765 YAG laser writer operating at 8 watts with 47.3 lines mm⁻¹ (1,200 line/inch) resolution. The resultant mask had a Dmin = 0.12 and Dmax = 3.05 (18A filter).

EXAMPLE 10:

[0113] The following formulation:

- (i) 0.50 grams 2,5-bis(4-diethylaminophenyl)-1,3,4-oxadiazole (Aldrich Chemical Co.), which is a fluorescent dye;
- (ii) 0.05 grams Cyasorb IR 165 (American Cyanamid);
- (iii) 6.00 grams topcoat lacquer (American Inks and Coatings Corp. product # 93-483); and
- (iv) 20.0 grams solvent (50% MEK, 20% MPK, 15% n-BuAc, 15% cyclohexanone),

was coated onto a 50% transmission aluminized polyester film as described in Example 1 with a # 2 Meyer rod and imagewise ablated onto an uncoated white bond paper using a Datrax 765 YAG laser writer operating at 7.5 watts with 47.3 lines mm⁻¹ (1,200 line/inch) resolution. The resultant image transferred onto the receptor sheet was barely visible in white light, but when viewed under ultraviolet light the image was highly visible. The imaged receptor sheet was not reproducible on a Minolta 7E7P5401 copier.

EXAMPLE 11:

[0114] The following samples of sputter coated metal films were overcoated with the cyan formulation described in Example 1 using a # 3 Meyer rod by hand drawdown. The actual thickness of the metal layer was unknown; accordingly, the films were characterized by their visible transmission densities (VTD) of the uncoated films. The films were imaged using a Nd:YAG laser at 1064 nm output delivering 0.10 J/cm², 0.16 J/cm² or 0.22 J/cm² in 104 nanoseconds using an uncoated newsprint receptor. The percentage of material transferred was calculated as in Example 4.

[0115] The results reported in the following Table IV were obtained:

TABLE IV

Metal VTD	Paper Dmax at J/cm ²			Imaged Film Dmin at 0.22 J/cm ²
	0.10	0.16	0.22	
Aluminum (vacuum metallized)				
0.77	30%	100%	100%	0.05
0.50	65%	100%	100%	0.05
0.43	95%	100%	100%	0.05
0.24	100%	100%	100%	0.05
0.15	80%	100%	100%	0.05
0.12	70%	100%	100%	0.05
0.10	45%	85%	100%	0.05
0.05	0%	10%	30%	0.05
Copper oxide (black)				
1.89	0%	100%	100%	0.12
0.98	0%	100%	100%	0.18
0.51	0%	75%	100%	0.15
Copper oxide (yellow)				
0.32	0%	0%	25%	0.65
0.19	0%	0%	35%	0.58

TABLE IV (continued)

Metal VTD Tin	Paper Dmax at J/cm ²			Imaged Film Dmin at 0.22 J/cm ²
0.4	35%	90%	100%	0.17
0.26	0%	55%	80%	0.17
0.12	0%	0%	30%	0.50
Indium/tin				
0.3	0%	65%	100%	0.13
0.19	0%	0%	35%	0.35
0.14	0%	0%	30%	0.44

[0116] In a separate experiment, the following dynamic release layers were compared at 0.16 J/cm²:

	Paper Dmax	Imaged Film Dmin
Aluminum (vacuum metallized) 0.30	100%	.05
Nickel/Chromium 0.24	100%	.04
Palladium 0.24	100%	.04

Claims

1. A method for transferring a contrasting pattern of intelligence from an ablation-transfer imaging medium (1, 2, 3; 2, 3, 7) to a receptor element (4) in contiguous registration therewith, the medium comprising:

- (i) a support substrate (1; 7)
- (ii) at least one intermediate dynamic release layer (2) essentially coextensive with the substrate (1; 7); and
- (iii) a radiation-ablative topcoat (3), also essentially coextensive with the substrate (1; 7),

the dynamic release layer and/or the topcoat containing at least one radiation-ablative binder,

the dynamic release layer (2) being capable of absorbing radiation (6) at a rate sufficient to effect the imagewise ablation mass transfer of the topcoat (3) to the receptor element (4), which method comprises imagewise irradiating the medium (1, 2, 3; 2, 3, 7) according to such pattern of intelligence with an intensity sufficient to effect the imagewise ablation mass transfer of the volume of the imagewise-exposed area of at least the topcoat (3) of the medium (1, 2, 3; 2, 3, 7) securely on to the receptor element (4),

characterized in that the dynamic release layer (2) comprises a non-black body radiation absorber but is free from black body radiation absorber, and in that the topcoat (3) comprises an imaging amount of a non-black body contrast imaging material, this contrast imaging material being transferred to the receptor element (4) to delineate the pattern of intelligence thereon.

2. A method according to claim 1 characterized in that the dynamic release layer (2) comprises a thin film of a metal, metal oxide or metal sulfide.

3. A method according to claim 2 characterized in that the dynamic release layer (2) comprises:

- a) a metal of Group Ib, IIb, IIIa, IVa, IVb, Va, Vb, VIa, VIb, VIIb or VIII of the Periodic Table;

- b) an alloy of the metals mentioned in paragraph a) above;
- c) an alloy of at least one of the metals mentioned in paragraph a) above with an element of Group Ia, IIa or IIIb of the Periodic Table;
- d) Al, Bi, Sn, In, or Zn, or an alloy thereof;
- e) a metal oxide or sulfide of Al, Bi, Sn, In, Zn, Ti, Cr, Mo, W, Co, Ir, Ni, Pd, Pt, Cu, Ag, Au, Zr or Te;
- f) Ge.

4. A method according to claim 1 characterized in that the dynamic release layer (2) comprises a sensitized liquid crystal material.
5. A method according to claim 1 characterized in that the dynamic release layer (2) comprises an organic material.
6. A method according to claim 5 characterized in that the organic material is:
 - a) a metal phthalocyanine, dithiolenes or anthraquinone compound; or
 - b) a polythiophene, polyaniline, polyacetylene, polyphenylene, polyphenylene sulfide, polypyrrole, or derivative or mixture thereof.
7. A method according to any one of the preceding claims characterized in that at least a portion of the dynamic release layer (2) is conjointly imagewise ablated on to the receptor element.
8. A method according to any one of the preceding claims characterized in that the dynamic release layer (2) has a thickness of up to 50nm (500 Angstroms).
9. A method according to any one of the preceding claims characterized in that the topcoat (3) comprises at least one radiation-ablative binder and at least one non-black body radiation absorber/sensitizer.
10. A method according to any one of the preceding claims characterized in that the contrast imaging material comprises a non-ablation sensitizing visible dye or pigment, ultraviolet or infrared absorbing material, magnetic material, polymeric material, fluorescent material, electrically conducting material, or mixture thereof.
11. A method according to any one of the preceding claims characterized in that the contrast imaging material is non-ablation sensitizing and highly absorbing in the spectral region of 350 nm to 450 nm.
12. A method according to claim 10 or 11 characterized in that the contrast imaging material comprises curcumin, an azo derivative, an oxadiazole derivative, a dicinnamalacetone derivative, a benzophenone derivative, a fluorescent oxazole, oxadiazole, coumarin or a carbostyryl derivative.
13. A method according to any one of the preceding claims characterized in that the substrate (1; 7) comprises glass, a polyester, a polycarbonate, a polyurethane, a polyolefin, a polyamide, a polysulfone, a polystyrene, a cellulosic material, aluminum or silicon.
14. A method according to any one of the preceding claims characterized in that the medium further comprises an anti-reflection layer (5) on the surface of the substrate (1; 7) opposite the topcoat (3).
15. A method according to any one of the preceding claims characterized in that the imagewise irradiation of the medium (1, 2, 3; 2, 3, 7) is effected with a Nd:YAG laser, a laser diode or an array of laser diodes.
16. A method according to any one of the preceding claims characterized in that the imagewise irradiation of the medium (1, 2, 3; 2, 3, 7) is effected at a power density of at least 10^4 watts/cm².
17. A method according to any one of the preceding claims characterized by being carried out successively employing a common receptor element but a plurality of ablation-transfer imaging media, the respective ablative topcoats of which including different contrast imaging colorants, and comprising transferring a multi-color proof to said common receptor element..
18. A method according to any one of the preceding claims characterized in that the dynamic release layer (2) comprises a laser radiation absorbing polymeric binder.

19. A method according to claim 9 characterized in that the radiation-ablative binder is decomposable by acid catalysis.

20. A method according to claim 19 characterized in that the radiation-ablative binder is adapted for acid catalyzed decomposition at a temperature of less than 200°C as measured under equilibrium conditions.

21. A method according to claim 9 characterized in that the radiation-ablative binder comprises a nitrocellulose, polycarbonate, polyurethane, polyester, polyorthoester or polyacetal.

22. A method according to claim 9 characterized in that the radiation-ablative binder comprises a polysulfone, styrene/acrylonitrile polymer, cellulosic ether or ester, polymethacrylate, polyvinylidene chloride, α -chloroacrylonitrile, maleic acid polymer, a copolymer or derivative thereof.

23. A method according to claim 9 characterized in that the radiation absorber/sensitizer comprises:

- a) an aromatic amine radical cation;
- b) a cation radical and an anion of a strong acid; or
- c) a cyanine or phthalocyanine dye, a metal dithiolene, a methylene blue salt, a di- or triarylmethane cation salt, a Wurster's blue salt or an onium salt.

24. A method according to claim 19 characterized in that the topcoat (3) further comprises at least one hydrogen atom donor for promoting acid formation for decomposition of the binder.

25. A method according to claim 24 characterized in that the hydrogen atom donor comprises an alcohol, thiol, phenol, amine or hydrocarbon.

26. An ablation-transfer imaging medium (1, 2, 3; 2, 3, 7) comprising:

- (i) a support substrate (1; 7)
- (ii) at least one intermediate dynamic release layer (2) essentially coextensive with the substrate (1; 7); and
- (iii) a radiation-ablative topcoat (3), also essentially coextensive with the substrate (1; 7),

the dynamic release layer and/or the topcoat containing at least one radiation-ablative binder, the dynamic release layer (2) being capable of absorbing radiation (6) at a rate sufficient to effect the imagewise ablation mass transfer of the topcoat (3) to the receptor element (4), characterized in that the dynamic release layer (2) comprises a non-black body radiation absorber but is free from black body radiation absorber, and in that the topcoat (3) comprises an imaging amount of a non-black body sensitizing contrast imaging material.

27. An ablation-transfer imaging medium according to claim 26 characterized in that the dynamic release layer (2) comprises a thin film of a metal, metal oxide or metal sulfide.

28. An ablation-transfer imaging medium according to claim 27 characterized in that the dynamic release layer (2) comprises:

- a) a metal of Group Ib, IIb, IIIa, IVa, IVb, Va, Vb, VIa, VIb, VIIb or VIII of the Periodic Table;
- b) an alloy of the metals mentioned in paragraph a) above;
- c) an alloy of at least one of the metals mentioned in paragraph a) above with an element of Group Ia, IIa or IIIb of the Periodic Table;
- d) Al, Bi, Sn, In or Zn, or an alloy thereof;
- e) a metal oxide or sulfide of Al, Bi, Sn, In, Zn, Ti, Cr, Mo, W, Co, Ir, Ni, Pd, Pt, Cu, Ag, Au, Zr or Te;
- f) Ge.

29. An ablation-transfer imaging medium according to claim 26 characterized in that the dynamic release layer (2) comprises a sensitized liquid crystal material.

30. An ablation-transfer imaging medium according to claim 26 characterized in that the dynamic release layer (2) comprises an organic material.

31. An ablation-transfer imaging medium according to claim 30 characterized in that the organic material is:

- a) a metal phthalocyanine, dithiolene or anthraquinone compound; or
- b) a polythiophene, polyaniline, polyacetylene, polyphenylene, polyphenylene sulfide, polypyrrole, or derivative or mixture thereof.

- 5 32. An ablation-transfer imaging medium according to any one of claims 26 to 31 characterized in that the dynamic release layer (2) has a thickness of up to 50nm (500 Angstroms).
33. An ablation-transfer imaging medium according to any one of claims 26 to 32 characterized in that the topcoat (3) comprises at least one radiation-ablative binder and at least one radiation absorber/sensitizer.
- 10 34. An ablation-transfer imaging medium according to any one of claims 26 to 33 characterized in that the contrast imaging material comprises a non-ablation sensitizing visible dye or pigment, ultraviolet or infrared absorbing material, magnetic material, polymeric material, fluorescent material, electrically conducting material, or mixture thereof.
- 15 35. An ablation-transfer imaging medium according to any one of claims 26 to 34 characterized in that the contrast imaging material is non-ablation sensitizing and highly absorbing in the spectral region of 350 nm to 450 nm.
- 20 36. An ablation-transfer imaging medium according to claim 34 or 35 characterized in that the contrast imaging material comprises curcumin, an azo derivative, an oxadiazole derivative, a dicinnamalacetone derivative, a benzophenone derivative, a fluorescent oxazole, oxadiazole, coumarin or a carbostyryl derivative.
- 25 37. An ablation-transfer imaging medium according to any one of claims 26 to 36 characterized in that the substrate (1; 7) comprises glass, a polyester, a polycarbonate, a polyurethane, a polyolefin, a polyamide, a polysulfone, a polystyrene, a cellulosic material, aluminum or silicon.
38. An ablation-transfer imaging medium according to any one of claims 26 to 37 characterized in that the medium further comprises an anti-reflection layer (5) on the surface of the substrate (1; 7) opposite the topcoat (3).
- 30 39. An ablation-transfer imaging medium according to any one of claims 26 to 37 characterized in that the dynamic release layer (2) comprises a laser radiation absorbing polymeric binder.
- 35 40. An ablation-transfer imaging medium according to claim 33 characterized in that the radiation-ablative binder is decomposable by acid catalysis.
- 40 41. An ablation-transfer imaging medium according to claim 40 characterized in that the radiation-ablative binder is adapted for acid catalyzed decomposition at a temperature of less than 200°C as measured under equilibrium conditions.
42. An ablation-transfer imaging medium according to claim 33 characterized in that the radiation-ablative binder comprises a nitrocellulose, polycarbonate, polyurethane, polyester, polyorthoester or polyacetal.
43. An ablation-transfer imaging medium according to claim 33 characterized in that the radiation-ablative binder comprises a polysulfone, styrene/acrylonitrile polymer, cellulosic ether or ester, polymethacrylate, polyvinylidene chloride, α -chloroacrylonitrile, maleic acid polymer, a copolymer or derivative thereof.
- 45 44. An ablation-transfer imaging medium according to claim 33 characterized in that the radiation absorber/sensitizer comprises:
 - 50 a) an aromatic amine radical cation;
 - b) a cation radical and an anion of a strong acid; or
 - c) a cyanine or phthalocyanine dye, a metal dithiolene, a methylene blue salt, a di- or triarylmethane cation salt, a Wurster's blue salt or an onium salt.
- 55 45. An ablation-transfer imaging medium according to claim 40 characterized in that the topcoat (3) further comprises at least one hydrogen atom donor for promoting acid formation for decomposition of the binder.
46. An ablation-transfer imaging medium according to claim 45 characterized in that the hydrogen atom donor com-

prises an alcohol, thiol, phenol, amine or hydrocarbon.

Patentansprüche

1. Verfahren zum Übertragen eines kontrastierenden Musters von Information von einem Ablationsübertragungs-Abbildungsmedium (1, 2, 3; 2, 3, 7) zu einem Rezeptor- bzw. Empfängerelement (4) in aneinandergrenzender Lagenaugkeit damit, wobei das Medium folgendes umfaßt:

(i) ein Trägersubstrat (1; 7)

(ii) wenigstens eine zwischenliegende dynamische Freisetzungs- bzw. Trennschicht (2), die im wesentlichen koextensiv mit dem Substrat (1; 7) ist bzw. sich im wesentlichen gemeinsam mit dem Substrat (1; 7) erstreckt; und

(iii) eine strahlungsablative obere Beschichtung bzw. dünne Oberschicht (3), die auch im wesentlichen koextensiv mit dem Substrat (1; 7) ist bzw. sich auch im wesentlichen gemeinsam mit dem Substrat (1; 7) erstreckt,

wobei die dynamische Freisetzungs- bzw. Trennschicht und/oder die obere Beschichtung bzw. dünne Oberschicht wenigstens ein strahlungsablatives Bindemittel enthält bzw. enthalten,

wobei die dynamische Freisetzungs- bzw. Trennschicht (2) fähig ist, Strahlung (6) mit einer Rate zu absorbieren, die genügend ist, um die bildweise Ablationsmassenübertragung der oberen Beschichtung bzw. dünnen Oberschicht (3) auf das Rezeptor- bzw. Empfängerelement (4) zu bewirken,

welches Verfahren das bildweise Bestrahlen des Mediums (1, 2, 3; 2, 3, 7) entsprechend einem solchen Muster von Information mit einer Intensität umfaßt, welche ausreicht, die bildweise Ablationsmassenübertragung des Volumens des bildweise exponierten Bereichs von wenigstens der oberen Beschichtung bzw. dünnen Oberschicht (3) des Mediums (1, 2, 3; 2, 3, 7) sicher auf das Rezeptor- bzw. Empfängerelement (4) zu bewirken,

dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) einen Nicht-Schwarzer-Körper-Strahlungsabsorber enthält, aber frei von Schwarzer-Körper-Strahlungsabsorber ist und daß die obere Beschichtung bzw. dünne Oberschicht (3) eine Abbildungsmenge von einem Nicht-Schwarzer-Körper-Kontrastabbildungsmaterial enthält, wobei dieses Kontrastabbildungsmaterial zu dem Rezeptor- bzw. Empfängerelement (4) übertragen wird, um das Muster an Information darauf darzustellen bzw. aufzuzeichnen.

2. Verfahren gemäß Anspruch 1, dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) einen dünnen Film aus einem Metall, Metall-oxid oder Metallsulfid umfaßt bzw. enthält.

3. Verfahren gemäß Anspruch 2, dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) folgendes umfaßt bzw. enthält:

a) ein Metall der Gruppe Ib, IIb, IIIa, IVa, IVb, Va, Vb, VIa, VIb, VIIb oder VIII des periodischen Systems;

b) eine Legierung der oben im Absatz a) erwähnten Metalle;

c) eine Legierung von wenigstens einem der im Absatz a) oben erwähnten Metalle mit einem Element der Gruppe Ia, IIa oder IIIb des periodischen Systems;

d) Al, Bi, Sn, In oder Zn, oder eine Legierung hiervon;

e) ein Metalloxid oder -sulfid von Al, Bi, Sn, In, Zn, Ti, Cr, Mo, W, Co, Ir, Ni, Pd, Pt, Cu, Ag, Au, Zr oder Te;

f) Ge.

4. Verfahren gemäß Anspruch 1, dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) ein sensibilisiertes Flüssigkristallmaterial umfaßt bzw. enthält.

5. Verfahren gemäß Anspruch 1, dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) ein organisches Material umfaßt bzw. enthält.
6. Verfahren gemäß Anspruch 5, dadurch **gekennzeichnet**, daß das organische Material folgendes ist:
 - a) eine Metallphthalocyanin-, Dithiolen- oder Anthrachinonverbindung; oder
 - b) ein Polythiophen, Polyanilin, Polyacetylen, Polyphenylen, Polyphenylensulfid, Polypyrrol, oder ein Abkömmling oder eine Mischung hiervon.
7. Verfahren gemäß irgendeinem der vorhergehenden Ansprüche, dadurch **gekennzeichnet**, daß wenigstens ein Teil der dynamischen Freisetzungs- bzw. Trennschicht (2) gemeinsam bildweise auf das Rezeptor- bzw. Empfängererelement ablatiert wird.
8. Verfahren gemäß irgendeinem der vorhergehenden Ansprüche, dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) eine Dicke von bis zu 50 nm (500 Å) hat.
9. Verfahren gemäß irgendeinem der vorhergehenden Ansprüche, dadurch **gekennzeichnet**, daß die obere Beschichtung bzw. dünne Oberschicht (3) wenigstens ein strahlungsablatives Bindemittel und wenigstens einen Nicht-Schwarzer-Körper-Strahlungsabsorber/Sensibilisierer umfaßt bzw. enthält.
10. Verfahren gemäß irgendeinem der vorhergehenden Ansprüche, dadurch **gekennzeichnet**, daß das Kontrastabbildungsmaterial einen Nichtablations-Sensibilisierungs-Sichtbar-Farbstoff oder ein Nichtablations-Sensibilisierungs-Sichtbar-Pigment, ein ultraviolett- oder infrarotabsorbierendes Material, ein magnetisches Material, ein Polymermaterial, ein fluoreszentes Material, ein elektrisch leitendes Material oder eine Mischung hiervon umfaßt bzw. enthält.
11. Verfahren gemäß irgendeinem der vorhergehenden Ansprüche, dadurch **gekennzeichnet**, daß das Kontrastabbildungsmaterial nichtablations-sensibilisierend und in hohem Maße absorbierend in dem Spektralbereich von 350 nm bis 450 nm ist.
12. Verfahren gemäß Anspruch 10 oder 11, dadurch **gekennzeichnet**, daß das Kontrastabbildungsmaterial Curcumin, ein Azo-Derivat, ein Oxadiazolderivat, ein Dicinnamalacetonderivat, ein Benzophenonderivat, ein fluoreszierendes Oxazol, Oxadiazol, Coumarin oder ein Carbostyrylderivat umfaßt bzw. enthält.
13. Verfahren gemäß irgendeinem der vorhergehenden Ansprüche, dadurch **gekennzeichnet**, daß das Substrat (1; 7)) Glas, ein Polyester, ein Polycarbonat, ein Polyurethan, ein Polyolefin, ein Polyamid, ein Polysulfon, ein Polystyrol, ein Cellulosematerial, Aluminium oder Silizium umfaßt bzw. enthält.
14. Verfahren gemäß irgendeinem der vorhergehenden Ansprüche, dadurch **gekennzeichnet**, daß das Medium weiter eine Antireflexionsschicht (5) auf der Oberfläche des Substrats (1; 7) gegenüber bzw. entgegengesetzt der obren Beschichtung bzw. dünnen Oberschicht (3) umfaßt bzw. enthält.
15. Verfahren gemäß irgendeinem der vorhergehenden Ansprüche, dadurch **gekennzeichnet**, daß die bildweise Bestrahlung des Mediums (1, 2, 3; 2, 3, 7) mit einem Nd:YAG-Laser, einer Laserdiode oder einer Anordnung bzw. Gruppierung von Laserdioden bewirkt wird.
16. Verfahren gemäß irgendeinem der vorhergehenden Ansprüche, dadurch **gekennzeichnet**, daß die bildweise Bestrahlung des Mediums (1, 2, 3; 2, 3, 7) mit einer Leistungsdichte von wenigstens 10^4 Watt/cm² bewirkt wird.
17. Verfahren gemäß irgendeinem der vorhergehenden Ansprüche, dadurch **gekennzeichnet**, daß es ausgeführt wird, indem aufeinanderfolgend ein gemeinsames Rezeptor- bzw. Empfängererelement, jedoch eine Mehrzahl von Ablationsübertragungsabbildungsmedien angewandt wird bzw. werden, wobei die jeweiligen ablativen oberen Beschichtungen bzw. dünnen Oberschichten hiervon unterschiedliche Kontrastabbildungsfärbemittel enthalten, und umfassend das Übertragen eines Multifarbproofs bzw. eines Multifarbprobedrucks auf das gemeinsame Rezeptor- bzw. Empfängererelement.
18. Verfahren gemäß irgendeinem der vorhergehenden Ansprüche, dadurch **gekennzeichnet**, daß die dynamische

Freisetzungs- bzw. Trennschicht (2) ein Laserstrahlung absorbierendes polymeres Bindemittel umfaßt bzw. enthält.

19. Verfahren gemäß Anspruch 9, dadurch **gekennzeichnet**, daß das strahlungsablative Bindemittel durch Säurekatalyse zersetzbar ist.

20. Verfahren gemäß Anspruch 19, dadurch **gekennzeichnet**, daß das strahlungsablative Bindemittel für eine säurekatalysierte Zersetzung bei einer Temperatur von weniger als 200°C, gemessen unter Gleichgewichtsbedingungen, geeignet ist.

21. Verfahren gemäß Anspruch 9, dadurch **gekennzeichnet**, daß das strahlungsablative Bindemittel Nitrocellulose, Polycarbonat, Polyurethan, Polyester, Polyorthoester oder Polyacetal umfaßt bzw. enthält.

22. Verfahren gemäß Anspruch 9, dadurch **gekennzeichnet**, daß das strahlungsablative Bindemittel ein Polysulfon, Styrol/Acrylnitril-Polymer, Celluloseether oder -ester, Polymethacrylat, Polyvinylidenchlorid, α -Chloroacrylnitril, Maleinsäurepolymer, ein Copolymer oder einen Abkömmling hiervon umfaßt bzw. enthält.

23. Verfahren gemäß Anspruch 9, dadurch **gekennzeichnet**, daß der Strahlungsabsorber/Sensibilisierer folgendes umfaßt bzw. enthält:

a) ein aromatisches Aminradikalkation;

b) ein Kationradikal und ein Anion einer starken Säure; oder

c) einen Zyanin- oder Phthalocyaninfarbstoff, ein Metaldithiolen, ein Methylenblausalz, ein Di- oder Triarylmethankationsalz, ein Wursters Blausalz oder ein Oniumsalz.

24. Verfahren gemäß Anspruch 19, dadurch **gekennzeichnet**, daß die obere Beschichtung bzw. dünne Oberschicht (3) weiter wenigstens einen Wasserstoffatomdonor zum Fördern der Säurebildung für die Zersetzung des Bindemittels umfaßt bzw. enthält.

25. Verfahren gemäß Anspruch 24, dadurch **gekennzeichnet**, daß der Wasserstoffatomdonor einen Alkohol, Thiol, Phenol, Amin oder Kohlenwasserstoff umfaßt bzw. enthält.

26. Ablationsübertragungsabbildungsmedium (1, 2, 3; 2, 3, 7), umfassend:

(i) ein Trägersubstrat (1; 7)

(ii) wenigstens eine zwischenliegende dynamische Freisetzungs- bzw. Trennschicht (2), die im wesentlichen koextensiv mit dem Substrat (1; 7) ist bzw. sich im wesentlichen gemeinsam mit dem Substrat (1; 7) erstreckt; und

(iii) eine strahlungsablative obere Beschichtung bzw. dünne Oberschicht (3), die auch im wesentlichen koextensiv mit dem Substrat (1; 7) ist bzw. sich auch im wesentlichen gemeinsam mit dem Substrat (1; 7) erstreckt,

wobei die dynamische Freisetzungs- bzw. Trennschicht und/oder die obere Beschichtung bzw. dünne Oberschicht wenigstens ein strahlungsablatives Bindemittel enthält bzw. enthalten,

wobei die dynamische Freisetzungs- bzw. Trennschicht (2) fähig ist, Strahlung (6) mit einer Rate zu absorbieren, die genügend ist, um die bildweise Ablationsmassenübertragung der oberen Beschichtung bzw. dünnen Oberschicht (3) auf das Rezeptor- bzw. Empfängerelement (4) zu bewirken, dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) einen Nicht-Schwarzer-Körper-Strahlungsabsorber umfaßt bzw. enthält, aber frei von Schwarzer-Körper-Strahlungsabsorber ist, und daß die obere Beschichtung bzw. dünne Oberschicht (3) eine Abbildungsmenge eines Nicht-Schwarzer-Körper-Sensibilisierungs-Kontrastabbildungsmaterial umfaßt bzw. enthält.

27. Ablationsübertragungsabbildungsmedium gemäß Anspruch 26, dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) einen dünnen Film aus einem Metall, Metall-oxid oder Metallsulfid umfaßt

bzw. enthält.

28. Ablationsübertragungsabbildungsmedium gemäß Anspruch 27, dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) folgendes umfaßt bzw. enthält:

a) ein Metall der Gruppe Ib, IIb, IIIa, IVa, IVb, Va, Vb, VIa, VIb, VIIb oder VIII des periodischen Systems;

b) eine Legierung der oben im Absatz a) erwähnten Metalle;

c) eine Legierung von wenigstens einem der oben im Absatz a) erwähnten Metalle mit einem Element der Gruppe Ia, IIa oder IIIb des periodischen Systems;

d) Al, Bi, Sn, In oder Zn, oder eine Legierung hiervon;

e) ein Metalloxid oder -sulfid von Al, Bi, Sn, In, Zn, Ti, Cr, Mo, W, Co, Ir, Ni, Pd, Pt, Cu, Ag, Au, Zr oder Te;

f) Ge.

29. Ablationsübertragungsabbildungsmedium gemäß Anspruch 26, dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) ein sensibilisiertes Flüssigkristallmaterial umfaßt bzw. enthält.

30. Ablationsübertragungsabbildungsmedium gemäß Anspruch 26, dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) ein organisches Material umfaßt bzw. enthält.

31. Ablationsübertragungsabbildungsmedium gemäß Anspruch 30, dadurch **gekennzeichnet**, daß das organische Material folgendes ist:

a) eine Metallphthalocyanin-, Dithiolen- oder Anthrachinonverbindung; oder

b) ein Polythiophen, Polyanilin, Polyacetylen, Polyphenylen, Polyphenylensulfid, Polypyrrol oder einen Abkömmling oder eine Mischung hiervon.

32. Ablationsübertragungsabbildungsmedium gemäß irgendeinem der Ansprüche 26 bis 31, dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) eine Dicke von bis zu 50 nm (500 Å) hat.

33. Ablationsübertragungsabbildungsmedium gemäß irgendeinem der Ansprüche 26 bis 32, dadurch **gekennzeichnet**, daß die obere Beschichtung bzw. dünne Oberschicht (3) wenigstens ein strahlungsablatives Bindemittel und wenigstens einen Strahlungsabsorber/Sensibilisierer umfaßt bzw. enthält.

34. Ablationsübertragungsabbildungsmedium gemäß irgendeinem der Ansprüche 26 bis 33, dadurch **gekennzeichnet**, daß das Konstrastabbildungsmaterial einen Nichtablations-Sensibilisierungs-Sichtbar-Farbstoff oder ein Nichtablations-Sensibilisierungs-Sichtbar-Pigment, ultraviolettoder infrarotabsorbierendes Material, magnetisches Material, Polymermaterial, fluoreszentes Material, elektrisch leitendes Material oder eine Mischung hiervon umfaßt bzw. enthält.

35. Ablationsübertragungsabbildungsmedium gemäß irgendeinem der Ansprüche 26 bis 34, dadurch **gekennzeichnet**, daß das Konstrastabbildungsmaterial nicht-ablationssensibilisierend und in hohem Maße absorbierend in dem Spektralbereich von 350 nm bis 450 nm ist.

36. Ablationsübertragungsabbildungsmedium gemäß Anspruch 34 oder 35, dadurch **gekennzeichnet**, daß das Konstrastabbildungsmaterial Curcumin, ein Azo-Derivat, ein Oxadiazolderivat, ein Dicinnamalacetonderivat, ein Benzophenonderivat, ein fluoreszierendes Oxazol, Oxadiazol, Coumarin oder ein Carbostyrylderivat umfaßt bzw. enthält.

37. Ablationsübertragungsabbildungsmedium gemäß irgendeinem der Ansprüche 26 bis 36, dadurch **gekennzeichnet**, daß das Substrat (1; 7) Glas, ein Polyester, ein Polycarbonat, ein Polyurethan, ein Polyolefin, ein Polyamid, ein Polysulfon, ein Polystyrol, ein Cellulosematerial, Aluminium oder Silizium umfaßt bzw. enthält.

38. Ablationsübertragungsabbildungsmedium gemäß irgendeinem der Ansprüche 26 bis 37, dadurch **gekennzeichnet**, daß das Medium weiter eine Antireflexionsschicht (5) auf der Oberfläche des Substrats (1; 7) gegenüber bzw. entgegengesetzt der oberen Beschichtung bzw. dünnen Oberschicht (3) umfaßt bzw. enthält.
- 5 39. Ablationsübertragungsabbildungsmedium gemäß irgendeinem der Ansprüche 26 bis 37, dadurch **gekennzeichnet**, daß die dynamische Freisetzungs- bzw. Trennschicht (2) ein Laserstrahlung absorbierendes Polymerbindemittel umfaßt bzw. enthält.
- 10 40. Ablationsübertragungsabbildungsmedium gemäß Anspruch 33, dadurch **gekennzeichnet**, daß das strahlungsablative Bindemittel durch Säurekatalyse zersetzbar ist.
- 15 41. Ablationsübertragungsabbildungsmedium gemäß Anspruch 40, dadurch **gekennzeichnet**, daß das strahlungsablative Bindemittel für eine säurekatalysierte Zersetzung bei einer Temperatur von weniger als 200°C, gemessen unter Gleichgewichtsbedingungen, geeignet ist.
- 20 42. Ablationsübertragungsabbildungsmedium gemäß Anspruch 33, dadurch **gekennzeichnet**, daß das strahlungsablative Bindemittel Nitrocellulose, Polycarbonat, Polyurethan, Polyester, Polyorthoester oder Polyacetal umfaßt bzw. enthält.
- 25 43. Ablationsübertragungsabbildungsmedium gemäß Anspruch 33, dadurch **gekennzeichnet**, daß das strahlungsablative Bindemittel ein Polysulfon, Styrol/Acrylnitril-Polymer, Celluloseether oder -ester, Polymethacrylat, Polyvinylidenchlorid, α -Chloroacrylnitril, Maleinsäurepolymer, ein Copolymer oder einen Abkömmling hiervon umfaßt bzw. enthält.
- 30 44. Ablationsübertragungsabbildungsmedium gemäß Anspruch 33, dadurch **gekennzeichnet**, daß der Strahlungsabsorber/Sensibilisierer folgendes umfaßt bzw. enthält:
- a) ein aromatisches Aminradikalkation;
 - b) ein Kationradikal und ein Anion einer starken Säure; oder
 - c) einen Cyanin- oder Phthalocyaninfarbstoff, ein Metalldithiolen, ein Methylenblausalz, ein Di- oder Triarylmethankationsalz, ein Wursters Blausalz oder ein Oniumsalz.
- 35 45. Ablationsübertragungsabbildungsmedium gemäß Anspruch 40, dadurch **gekennzeichnet**, daß die obere Beschichtung bzw. dünne Oberschicht (3) weiter wenigstens einen Wasserstoffatomdonor zum Fördern der Säurebildung für die Zersetzung des Bindemittels umfaßt bzw. enthält.
- 40 46. Ablationsübertragungsabbildungsmedium gemäß Anspruch 45, dadurch **gekennzeichnet**, daß der Wasserstoffatomdonor einen Alkohol, Thiol, Phenol, Amin oder Kohlenwasserstoff umfaßt bzw. enthält.

Revendications

- 45 1. Procédé de transfert d'un motif contrasté formant informations à partir d'un support d'imagerie à transfert par ablation (1, 2, 3 ; 2, 3, 7) vers un élément récepteur (4) selon un repérage contigu avec ce dernier, le support comprenant :
- 50 (i) un substrat de support (1 ; 7)
 (ii) au moins une couche de libération dynamique intermédiaire (2) sensiblement co-extensive avec le substrat (1 ; 7) ; et
 (iii) une couche de finition ablative par rayonnement (3), également sensiblement co-extensive avec le substrat (1 ; 7),
- 55 la couche de libération dynamique et/ou la couche de finition contenant au moins un liant ablatif par rayonnement,
- la couche de libération dynamique (2) étant susceptible d'absorber un rayonnement (6) à une vitesse suffisante

pour effectuer le transfert de matière par ablation, en ce qui concerne l'image, de la couche de finition (3) vers l'élément récepteur (4),

lequel procédé comprend l'irradiation, en ce qui concerne l'image, du support (1, 2, 3 ; 2, 3, 7) selon un motif formant informations de ce type avec une intensité suffisante pour effectuer le transfert de matière par ablation, en ce qui concerne l'image, du volume de la zone exposée concernant l'image d'au moins la couche de finition (3) du support (1, 2, 3 ; 2, 3, 7) de manière sûre sur l'élément récepteur (4),

caractérisé en ce que la couche de libération dynamique (2) comprend un absorbeur de rayonnement formant corps non noir mais est exempte d'un absorbeur de rayonnement formant corps noir, et en ce que la couche de finition (3) comprend une quantité d'imagerie d'une matière d'imagerie de contraste formant corps non noir, cette matière d'imagerie de contraste étant transférée sur l'élément récepteur (4) pour délimiter le motif formant informations sur ce dernier.

2. Procédé selon la revendication 1, caractérisé en ce que la couche de libération dynamique (2) comprend un film mince de métal, d'oxyde de métal ou de sulfure de métal.

3. Procédé selon la revendication 2, caractérisé en ce que la couche de libération dynamique (2) comprend :

- a) un métal du groupe Ib, IIb, IIIa, IVa, IVb, Va, Vb, Via, VIb, VIIb ou VIII de la table périodique des éléments ;
- b) un alliage des métaux mentionnés au paragraphe a) précédent ;
- c) un alliage d'au moins un des métaux mentionnés au paragraphe a) précédent avec un élément du groupe Ia, IIa ou IIIb de la table périodique des éléments ;
- d) Al, Bi, Sn, In ou Zn, ou un alliage de ces derniers ;
- e) un oxyde de métal ou un sulfure de Al, Bi, Sn, In, Zn, Ti, Cr, Mo, W, Co, Ir, Ni, Pd, Pt, Cu, Ag, Au, Zr ou Te ;
- f) Ge.

4. Procédé selon la revendication 1, caractérisé en ce que la couche de libération dynamique (2) comprend une matière sensible à cristaux liquides.

5. Procédé selon la revendication 1, caractérisé en ce que la couche de libération dynamique (2) comprend une matière organique.

6. Procédé selon la revendication 5, caractérisé en ce que la matière organique est :

- a) un composé métallique de phtalocyanine, de dithiolène ou d'anthraquinone ; ou
- b) du polythiophène, de la polyaniline, du polyacétylène, du polyphénylène, du sulfure de polyphénylène, du polypyrrole, ou un dérivé ou un mélange de ces derniers.

7. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce qu'au moins une partie de la couche de libération dynamique (2) subit une ablation de manière conjointe, en ce qui concerne l'image, sur l'élément récepteur.

8. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que la couche de libération dynamique (2) a une épaisseur allant jusqu'à 50 nm (500 Angströms).

9. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que la couche de finition (3) comprend au moins un liant ablatif par rayonnement et au moins un absorbeur/sensibilisateur de rayonnement formant corps non noir.

10. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que la matière d'imagerie contrastée comprend un colorant ou un pigment visible sensible non ablatif, une matière absorbant les ultraviolets ou l'infrarouge, une matière magnétique, une matière polymère, une matière fluorescente, une matière électriquement conductrice, ou un mélange de ces dernières.

11. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que la matière d'imagerie contrastée n'est pas sensible à l'ablation et est hautement absorbante dans la zone spectrale de 350 nm à 450 nm.

12. Procédé selon la revendication 10 ou 11, caractérisé en ce que la matière d'imagerie contrastée comprend du

jaune de curcuma, un dérivé azoïque, un dérivé d'oxadiazole, un dérivé d'acétone cinnamique, un dérivé de benzophénone, un oxazole fluorescent, un oxadiazole, de la coumarine ou un dérivé de carbostyryle.

13. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que le substrat (1 ; 7) comprend du verre, un polyester, un polycarbonate, un polyuréthane, une polyoléfine, un polyamide, un polysulfone, un polystyrène, une matière cellulosique, de l'aluminium ou du silicium.
14. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que le support comprend de plus une couche antiréfléchissante (5) sur la surface du substrat (1 ; 7) opposée à la couche de finition (3).
15. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que l'irradiation, en ce qui concerne l'image, du support (1, 2, 3 ; 2, 3, 7) est effectuée avec un laser YAG au néodyme, une diode laser ou un groupement de diodes laser.
16. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que l'irradiation, en ce qui concerne l'image, du support (1, 2, 3 ; 2, 3, 7) est effectuée à une densité de puissance d'au moins 10^4 watts/cm².
17. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce qu'il est exécuté successivement en utilisant un élément récepteur commun mais une pluralité de supports d'imagerie à transfert par ablation, dont les couches de finition ablatives respectives comprennent différents colorants d'imagerie contrastés, et en ce qu'il comprend le transfert d'une épreuve multicolore sur ledit élément récepteur commun.
18. Procédé selon l'une quelconque des revendications précédentes, caractérisé en ce que la couche de libération dynamique (2) comprend un liant polymère absorbant le rayonnement laser.
19. Procédé selon la revendication 9, caractérisé en ce que le liant ablatif par rayonnement peut être décomposé par une catalyse d'acide.
20. Procédé selon la revendication 19, caractérisé en ce que le liant ablatif par rayonnement est conçu pour une décomposition catalysée d'acide à une température de moins de 200 °C lorsque mesurée sous des conditions d'équilibre.
21. Procédé selon la revendication 9, caractérisé en ce que le liant ablatif par rayonnement comprend de la nitrocellulose, du polycarbonate, du polyuréthane, du polyester, du polyortho-ester ou du polyacétal.
22. Procédé selon la revendication 9, caractérisé en ce que le liant ablatif par rayonnement comprend un polysulfone, un polymère de styrène/acrylonitrile, un éther ou un ester cellulosique, du polyméthacrylate, du chlorure de polyvinylidène, un α -chloro-acrylonitrile, un polymère d'acide maléique, un copolymère ou un dérivé de ces derniers.
23. Procédé selon la revendication 9, caractérisé en ce que l'absorbeur/sensibilisateur de rayonnement comprend :
 - a) un cation à radical amine aromatique ;
 - b) un radical cationique et un anion d'un acide fort ; ou
 - c) un colorant de cyanine ou de phthalocyanine, un dithiolène métallique, un sel de bleu de méthylène, un sel cationique de di- ou de triarylméthane, un sel de bleu de Wurster ou un sel d'onium.
24. Procédé selon la revendication 19, caractérisé en ce que la couche de finition (3) comprend au moins un atome d'hydrogène donneur pour favoriser la formation d'acide pour la décomposition du liant.
25. Procédé selon la revendication 24, caractérisé en ce que l'atome d'hydrogène donneur comprend un alcool, un thiol, un phénol, une amine ou un hydrocarbure.
26. Support d'imagerie à transfert par ablation (1, 2, 3 ; 2, 3, 7) comprenant :
 - (i) un substrat de support (1 ; 7)
 - (ii) au moins une couche de libération dynamique intermédiaire (2) sensiblement co-extensive avec le substrat (1 ; 7) ; et
 - (iii) une couche de finition ablatif par rayonnement (3), également sensiblement co-extensive avec le substrat

(1 ; 7),

la couche de libération dynamique et/ou la couche de finition contenant au moins un liant ablatif par rayonnement,

la couche de libération dynamique (2) étant susceptible d'absorber un rayonnement (6) à une vitesse suffisante pour effectuer le transfert de matière par ablation, en ce qui concerne l'image, de la couche de finition (3) vers l'élément récepteur (4),

caractérisé en ce que la couche de libération dynamique (2) comprend un absorbeur de rayonnement formant corps non noir mais est exempte d'un absorbeur de rayonnement formant corps noir, et en ce que la couche de finition (3) comprend une quantité d'imagerie d'une matière d'imagerie de contraste formant corps non noir.

27. Support d'imagerie à transfert par ablation selon la revendication 26, caractérisé en ce que la couche de libération dynamique (2) comprend un film mince de métal, d'oxyde de métal ou de sulfure de métal.

28. Support d'imagerie à transfert par ablation selon la revendication 27, caractérisé en ce que la couche de libération dynamique (2) comprend :

a) un métal du groupe Ib, IIb, IIIa, IVa, IVb, Va, Vb, VIa, VIb, VIIb ou VIII de la table périodique des éléments ;

b) un alliage des métaux mentionnés au paragraphe a) précédent ;

c) un alliage d'au moins un des métaux mentionnés au paragraphe a) précédent avec un élément du groupe Ia, IIa ou IIIb de la table périodique des éléments ;

d) Al, Bi, Sn, In ou Zn, ou un alliage de ces derniers ;

e) un oxyde de métal ou un sulfure de Al, Bi, Sn, In, Zn, Ti, Cr, Mo, W, Co, Ir, Ni, Pd, Pt, Cu, Ag, Au, Zr ou Te ;

f) Ge.

29. Support d'imagerie à transfert par ablation selon la revendication 26, caractérisé en ce que la couche de libération dynamique (2) comprend une matière sensible à cristaux liquides.

30. Support d'imagerie à transfert par ablation selon la revendication 26, caractérisé en ce que la couche de libération dynamique (2) comprend une matière organique.

31. Support d'imagerie à transfert par ablation selon la revendication 30, caractérisé en ce que la matière organique est :

a) un composé métallique de phtalocyanine, de dithiolène ou d'anthraquinone ; ou

b) du polythiophène, de la polyaniline, du polyacétylène, du polyphénylène, du sulfure de polyphénylène, du polypyrrole, ou un dérivé ou un mélange de ces derniers.

32. Support d'imagerie à transfert par ablation selon l'une quelconque des revendications 26 à 31, caractérisé en ce que la couche de libération dynamique (2) a une épaisseur allant jusqu'à 50 nm (500 Angströms).

33. Support d'imagerie à transfert par ablation selon l'une quelconque des revendications 26 à 32, caractérisé en ce que la couche de finition (3) comprend au moins un liant ablatif par rayonnement et au moins un absorbeur/sensibilisateur de rayonnement.

34. Support d'imagerie à transfert par ablation selon l'une quelconque des revendications 26 à 33, caractérisé en ce que la matière d'imagerie contrastée comprend un colorant ou un pigment visible sensible non ablatif, une matière absorbant les ultraviolets ou l'infrarouge, une matière magnétique, une matière polymère, une matière fluorescente, une matière électriquement conductrice, ou un mélange de ces dernières.

35. Support d'imagerie à transfert par ablation selon l'une quelconque des revendications 26 à 34, caractérisé en ce que la matière d'imagerie contrastée n'est pas sensible à l'ablation et est hautement absorbante dans la zone spectrale de 350 nm à 450 nm.

36. Support d'imagerie à transfert par ablation selon la revendication 34 ou 35, caractérisé en ce que la matière d'imagerie contrastée comprend du jaune de curcuma, un dérivé azoïque, un dérivé d'oxadiazole, un dérivé d'acétone dicinnamique, un dérivé de benzophénone, un oxazole fluorescent, un oxadiazole, de la coumarine ou un dérivé de carbostyryle.

37. Support d'imagerie à transfert par ablation selon l'une quelconque des revendications 26 à 36, caractérisé en ce que le substrat (1 ; 7) comprend du verre, un polyester, un polycarbonate, un polyuréthane, une polyoléfine, un polyamide, un polysulfone, un polystyrène, une matière cellulosique, de l'aluminium ou du silicium.

38. Support d'imagerie à transfert par ablation selon l'une quelconque des revendications 26 à 37, caractérisé en ce que le support comprend de plus une couche antiréfléchissante (5) sur la surface du substrat (1 ; 7) opposée à la couche de finition (3).

39. Support d'imagerie à transfert par ablation selon l'une quelconque des revendications 26 à 37, caractérisé en ce que la couche de libération dynamique (2) comprend un liant polymère absorbant le rayonnement laser.

40. Support d'imagerie à transfert par ablation selon la revendication 33, caractérisé en ce que le liant ablatif par rayonnement peut être décomposé par une catalyse d'acide.

41. Support d'imagerie à transfert par ablation selon la revendication 40, caractérisé en ce que le liant ablatif par rayonnement est conçu pour une décomposition catalysée d'acide à une température de moins de 200 °C lorsque mesurée sous des conditions d'équilibre.

42. Support d'imagerie à transfert par ablation selon la revendication 33, caractérisé en ce que le liant ablatif par rayonnement comprend de la nitrocellulose, du polycarbonate, du polyuréthane, du polyester, du polyortho-ester ou du polyacétal.

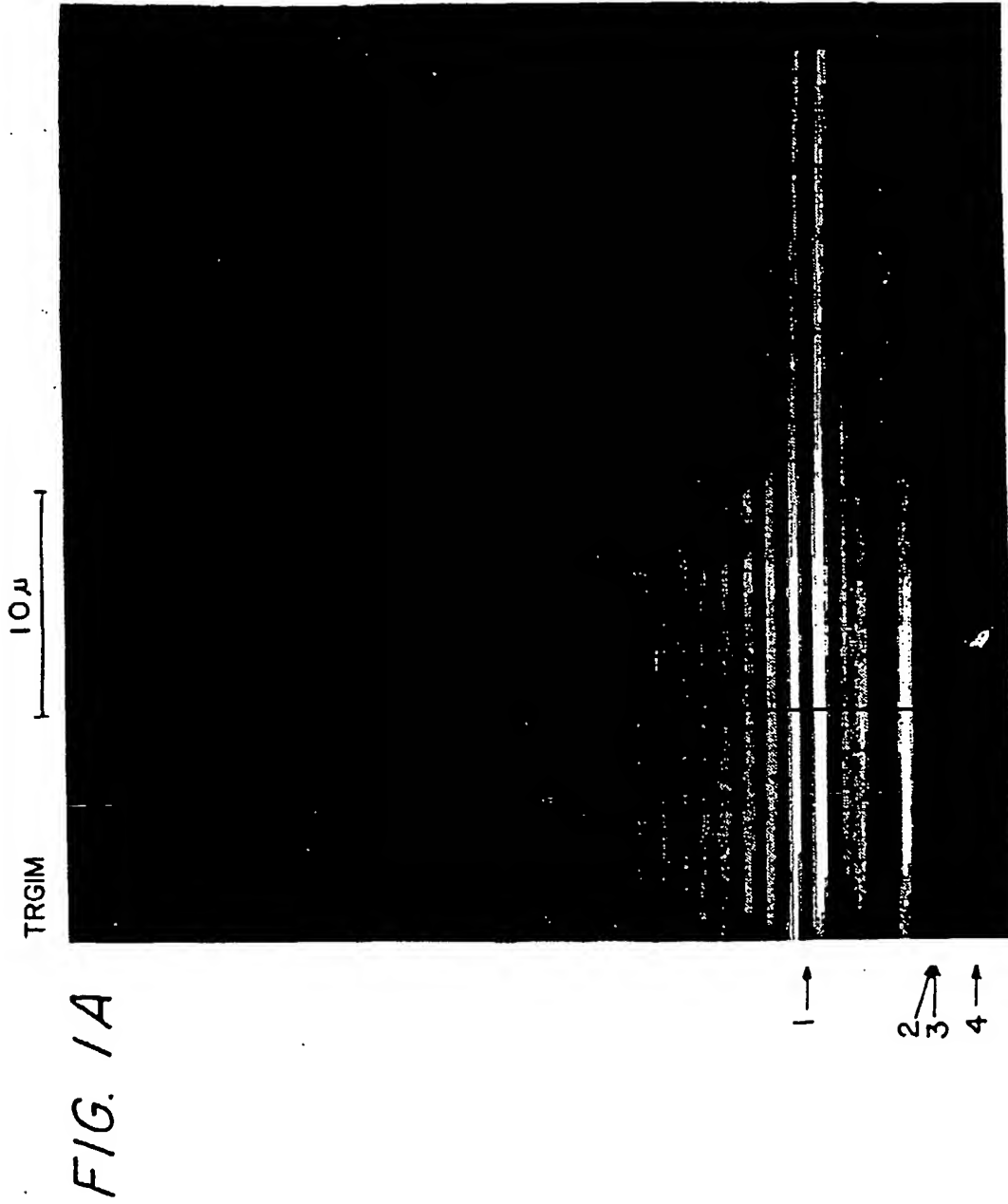
43. Support d'imagerie à transfert par ablation selon la revendication 33, caractérisé en ce que le liant ablatif par rayonnement comprend un polysulfone, un polymère de styrène/acrylnitrile, un éther ou un ester cellulosique, du polyméthacrylate, du chlorure de polyvinylidène, un α -chloro-acrylnitrile, un polymère d'acide maléique, un copolymère ou un dérivé de ces derniers.

44. Support d'imagerie à transfert par ablation selon la revendication 33, caractérisé en ce que l'absorbeur/sensibilisateur de rayonnement comprend :

- a) un cation à radical amine aromatique ;
- b) un radical cationique et un anion d'un acide fort ; ou
- c) un colorant de cyanine ou de phthalocyanine, un dithiolène métallique, un sel de bleu de méthylène, un sel cationique de di- ou de triarylméthane, un sel de bleu de Wurster ou un sel d'onium.

45. Support d'imagerie à transfert par ablation selon la revendication 40, caractérisé en ce que la couche de finition (3) comprend au moins un atome d'hydrogène donneur pour favoriser la formation d'acide pour la décomposition du liant.

46. Support d'imagerie à transfert par ablation selon la revendication 45, caractérisé en ce que l'atome d'hydrogène donneur comprend un alcool, un thiol, un phénol, une amine ou un hydrocarbure.



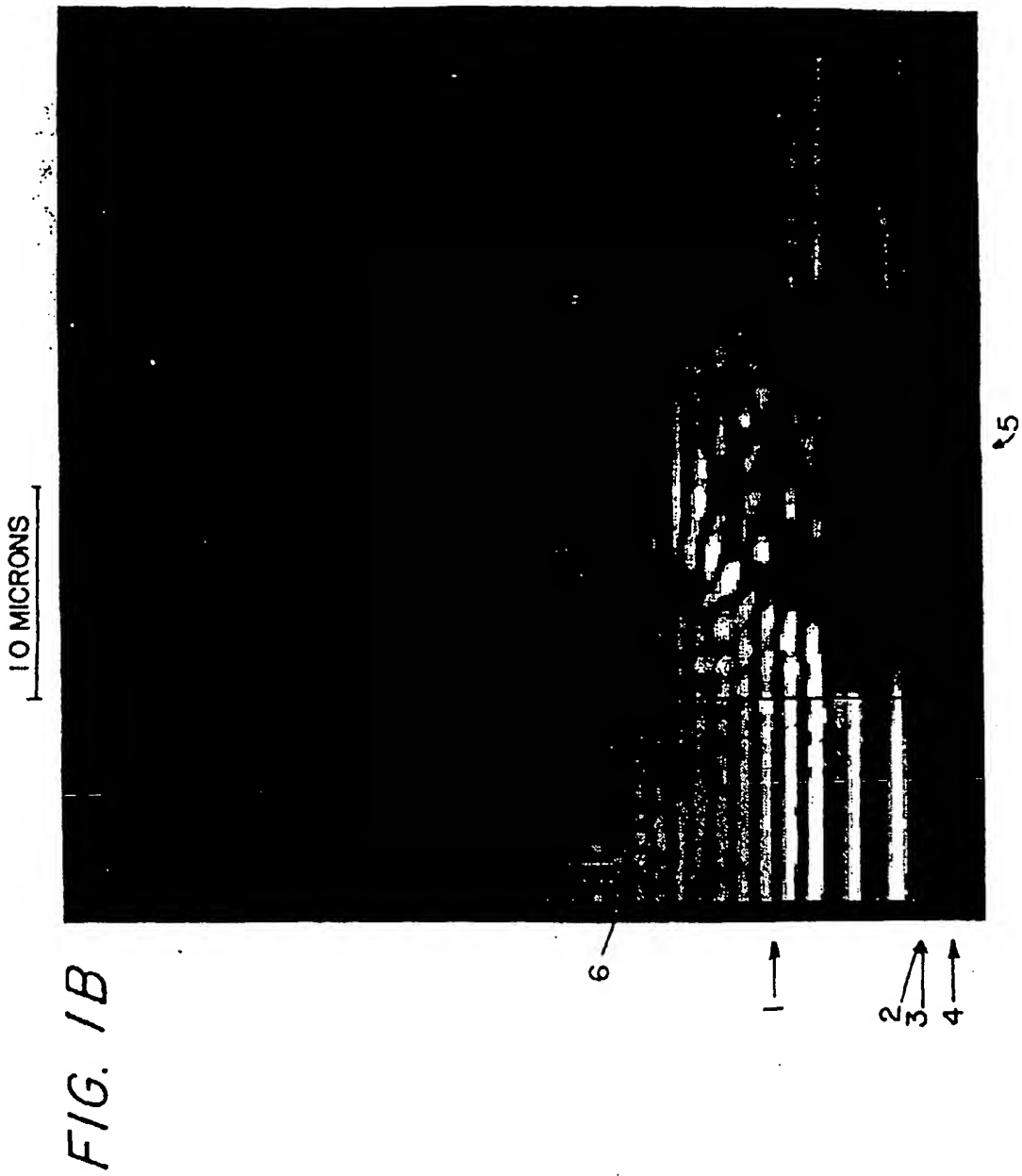


FIG. 2A

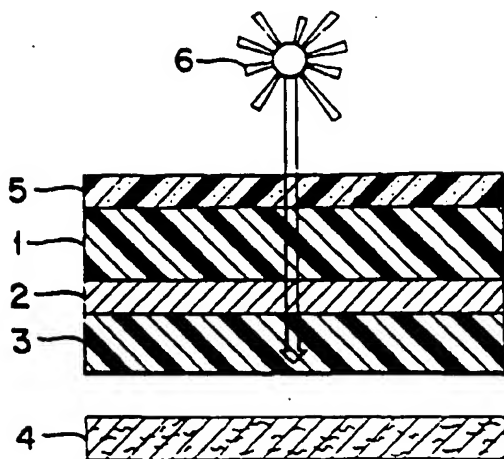


FIG. 2B

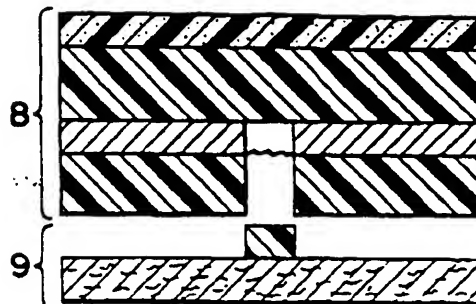


FIG. 3A

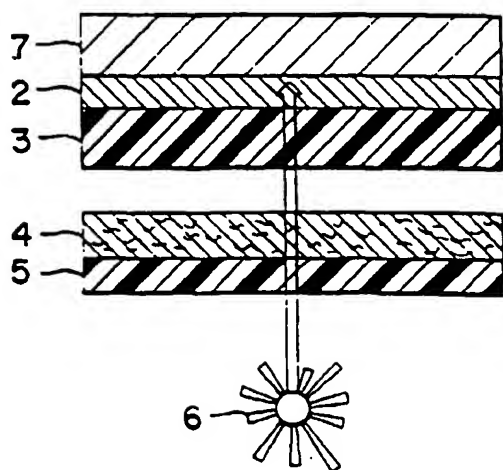
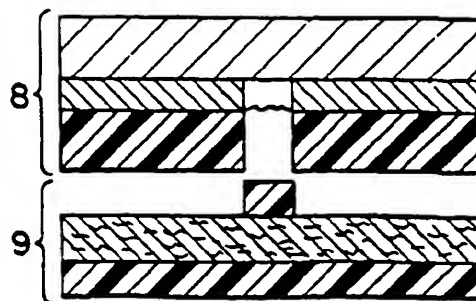
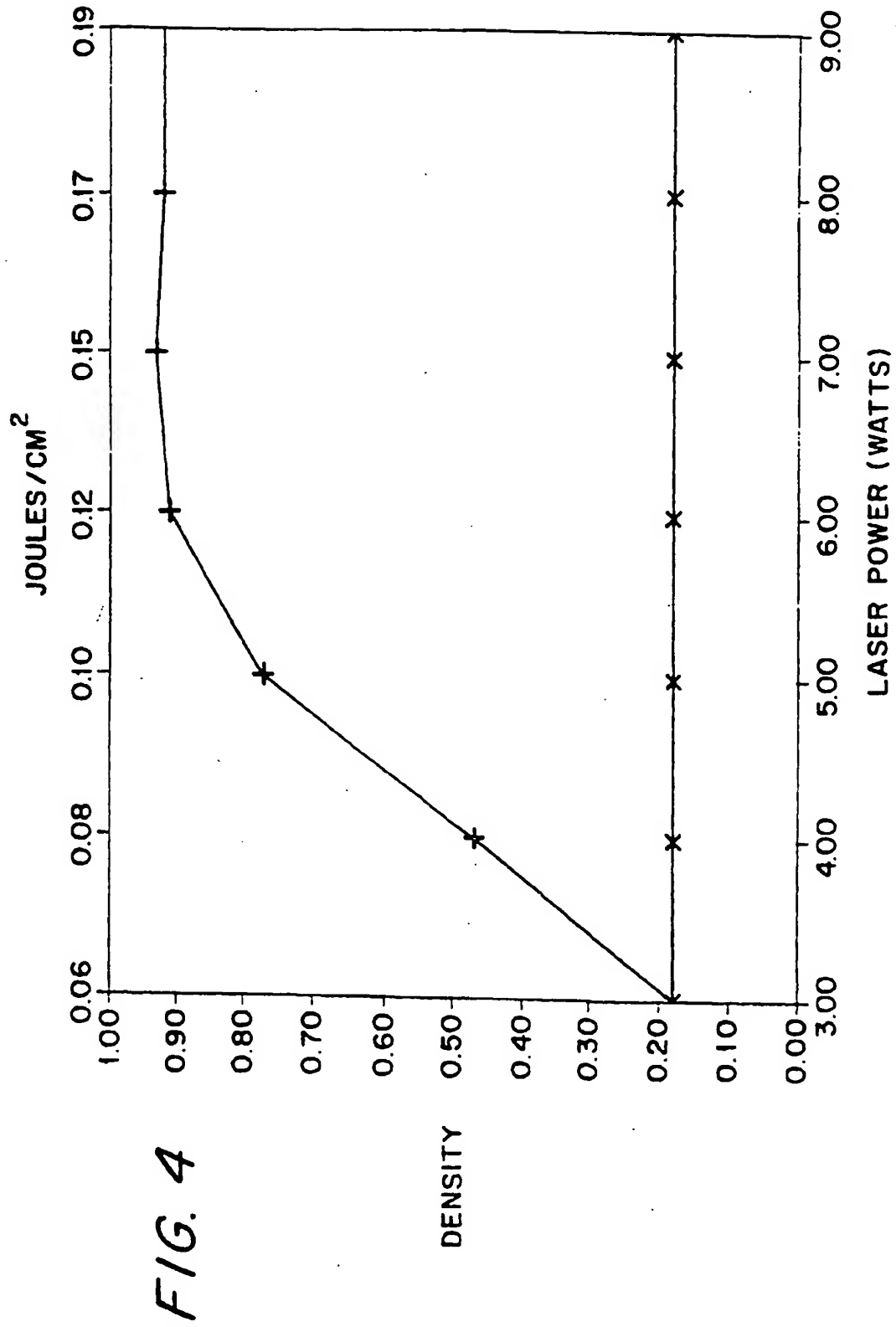


FIG. 3B





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